

=&gt; d que

L1 16 SEA FILE=REGISTRY ABB=ON PLU=ON RRRPRPPYLPRRPP/SQSP  
 L2 2 SEA FILE=REGISTRY ABB=ON PLU=ON ("PR 39"/CN OR "PR 39 (ION  
 EXCHANGER)"/CN OR "PR 39 (PEPTIDE)"/CN)  
 L4 53 SEA FILE=HCAPLUS ABB=ON PLU=ON L1 AND (L2 OR PR39 OR PR 39)  
 L5 8645 SEA FILE=HCAPLUS ABB=ON PLU=ON ANGIOGENESIS+NT/CT  
 L6 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L1 AND (L5 OR ANGIOGEN?)  
 L7 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L4 AND (L5 OR ANGIOGEN?)  
 L8 6 SEA FILE=HCAPLUS ABB=ON PLU=ON L6 OR L7

=&gt; d ibib abs hitstr l8 1-6

L8 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:220416 HCAPLUS

DOCUMENT NUMBER: 136:257252

TITLE: Method of modulating neovascularization

INVENTOR(S): Kovesdi, Imre

PATENT ASSIGNEE(S): Genvec, Inc., USA

SOURCE: PCT Int. Appl., 33 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002022176	A1	20020321	WO 2001-US28954	20010914
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2001091019	A5	20020326	AU 2001-91019	20010914
PRIORITY APPLN. INFO.: US 2000-233001P P 20000915				
WO 2001-US28954 W 20010914				
AB	The present invention provides a method of modulating neovascularization in an animal. The method comprises administering to the animal two or more nucleic acid sequences, each nucleic acid sequence encoding at least one <b>angiogenesis</b> -modulation factor that acts upon a different <b>angiogenic</b> process, such that the nucleic acid sequences are expressed to produce the <b>angiogenesis</b> -modulation factors to modulate neovascularization in the animal. Modulating neovascularization includes the induction of neovascularization or, in the alternative, the inhibition or redn. of neovascularization.			
IT	139637-11-9, PR39 peptide			
	RL: BUU (Biological use, unclassified); PAC (Pharmacological activity); BIOL (Biological study); USES (Uses)			
	(method of modulating neovascularization)			
RN	139637-11-9 HCAPLUS			
CN	L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-			

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prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:319740 HCAPLUS

DOCUMENT NUMBER: 134:336214

TITLE: Method for **PR-39** peptide regulated stimulation of **angiogenesis**

INVENTOR(S): Simons, Michael; Gao, Youhe

PATENT ASSIGNEE(S): Beth Israel Deaconess Medical Center, USA

SOURCE: PCT Int. Appl., 52 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001030368	A1	20010503	WO 2000-US27552	20001006
W: AU, CA, JP				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

PRIORITY APPLN. INFO.: US 1999-426011 A 19991025

AB The present invention provides both a method and means for regulating **angiogenesis** within living cells, tissues, and organs in-situ. The regulation is performed using native **PR-39** peptide or one of its shorter-length homolog, for interaction with such proteasomes as one present in the cytoplasm of viable cells. The result of **PR-39** peptide interaction with proteasomes is a decrease in the intracellular degrdn. of active peptides such as HIF-1.alpha. and a consequential stimulation of **angiogenesis** in-situ.

IT 298702-64-4

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)  
(**PR-39** peptide regulated stimulation of **angiogenesis**)

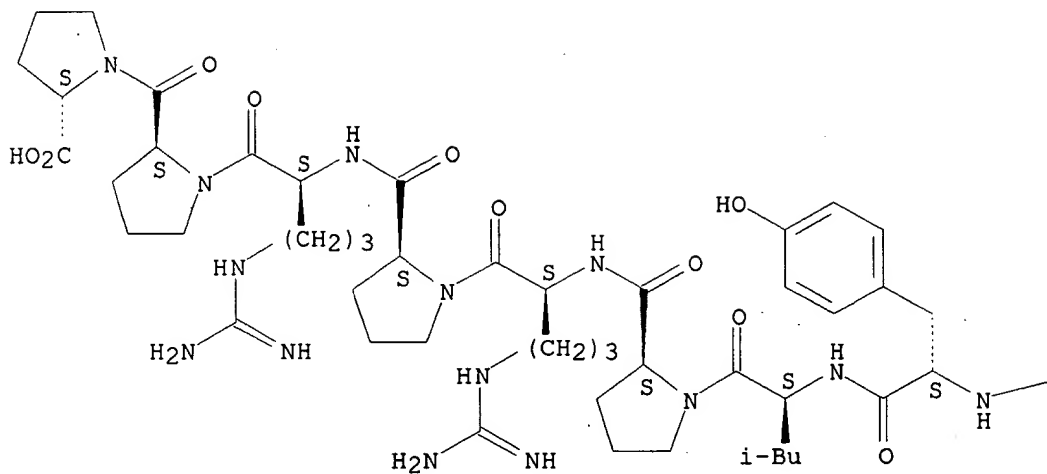
RN 298702-64-4 HCAPLUS

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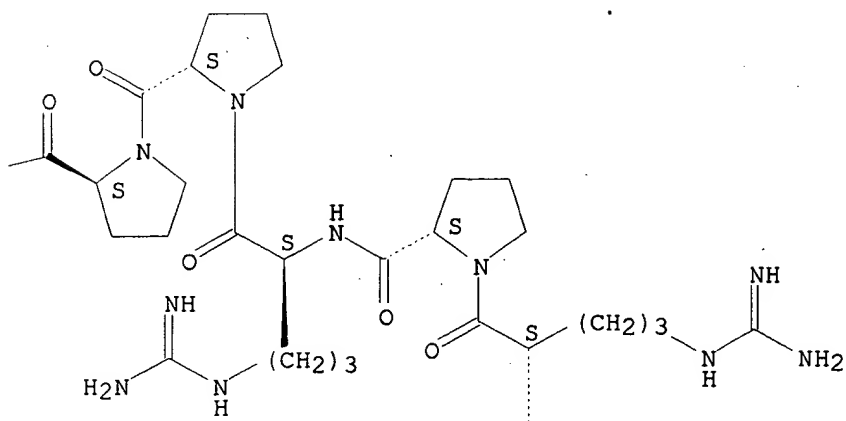
Absolute stereochemistry.

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PAGE 1-A

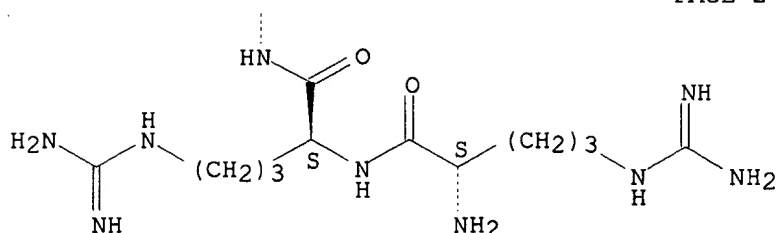


PAGE 1-B



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PAGE 2-B



IT 139637-11-9, PR-39 peptide

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(PR-39 peptide regulated stimulation of angiogenesis)

RN 139637-11-9 HCAPLUS

CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2000:706997 HCAPLUS

DOCUMENT NUMBER: 133:276343

TITLE: Method for PR-39 peptide regulated stimulation of angiogenesis

INVENTOR(S): . Simons, Michael; Gao, Youhe

PATENT ASSIGNEE(S): Beth Israel Deaconess Medical Center, USA

SOURCE: PCT Int. Appl., 51 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000057895	A1	20001005	WO 2000-US7050	20000316

W: AU, CA, JP

RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE

EP 1165111      A1      20020102      EP 2000-919442      20000316

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
IE, FI

PRIORITY APPLN. INFO.: US 1999-276868 A 19990326

WO 2000-US7050 W 200000316

AB The present invention provides both a method and means for regulating **angiogenesis** within living cells, tissues, and organs in-situ. The regulation is performed using native **PR-39** peptide

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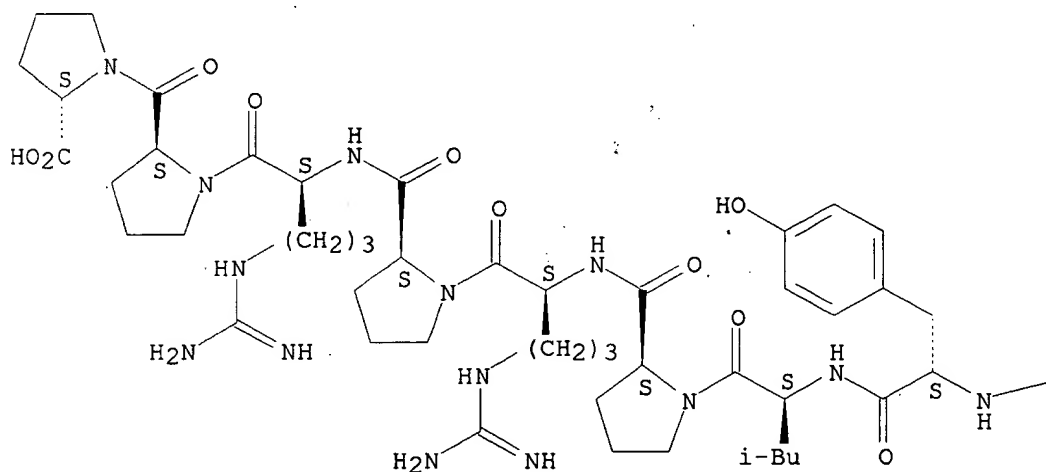


IT 298702-64-4P

(PR-39 peptide-regulated stimulation of angiogenesis)

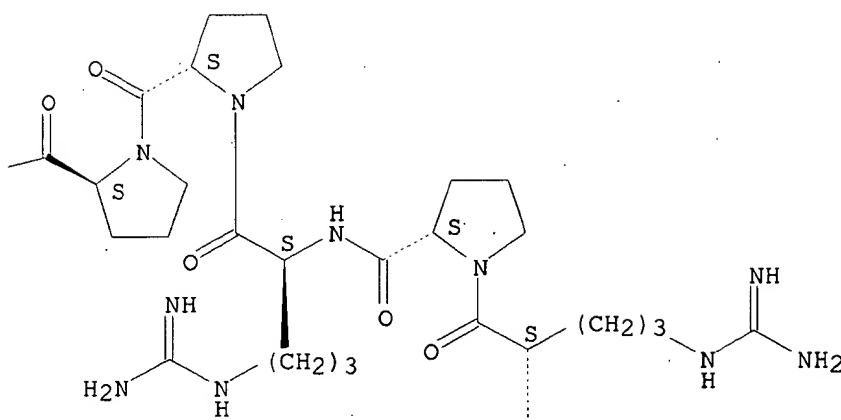
CN L-Proline, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-  
(9CI) (CA INDEX NAME)

PAGE 1-A

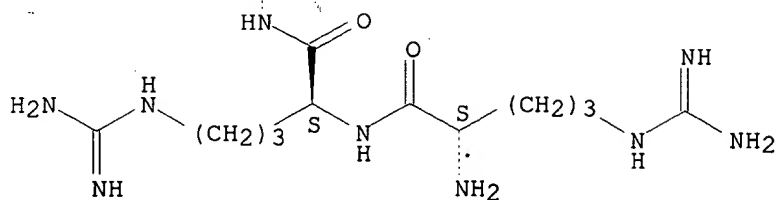


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PAGE 1-B



PAGE 2-B



IT 148046-54-2

RL: PRP (Properties)

(unclaimed protein sequence; method for PR-39  
peptide regulated stimulation of **angiogenesis**)

RN 148046-54-2 HCAPLUS

CN L-Proline, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 4 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2000:178321 HCAPLUS

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DOCUMENT NUMBER: 133:205925  
TITLE: **PR39**, a peptide regulator of **angiogenesis**. [Erratum to document cited in CA132:149677]  
AUTHOR(S): Li, Jian; Post, Mark; Volk, Rudiger; Gao, Youhe; Li, Min; Metals, Caroline; Sato, Kaori; Tsai, Jo; Aird, William; Rosenberg, Robert D.; Hampton, Thomas G.; Li, Jianyi; Sellke, Frank; Carmeliet, Peter; Simons, Michael  
CORPORATE SOURCE: Angiogenesis Research Center, Department of Surgery, Beth Israel Deaconess Medical Center and Harvard Medical School, Boston, MA, 02215, USA  
SOURCE: Nature Medicine (New York) (2000), 6(3), 356  
CODEN: NAMEFI; ISSN: 1078-8956  
PUBLISHER: Nature America  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
AB The correct versions are given for Figs. 2a, c, and d on page 51; Fig. 3c on page 52; and Fig. 5b on page 53.  
IT **139637-11-9, PR-39**  
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(**PR39** peptide in regulation of **angiogenesis** by inhibiting ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein (Erratum))  
RN 139637-11-9 HCAPLUS  
CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

L8 ANSWER 5 OF 6 HCAPLUS COPYRIGHT 2003 ACS  
ACCESSION NUMBER: 2000:46162 HCAPLUS  
DOCUMENT NUMBER: 132:149677  
TITLE: **PR39**, a peptide regulator of **angiogenesis**  
AUTHOR(S): Li, Jian; Post, Mark; Volk, Rudiger; Gao, Youhe; Li, Min; Metals, Caroline; Sato, Kaori; Tsai, Jo; Aird, William; Rosenberg, Robert D.; Hampton, Thomas G.; Li, Jianyi; Sellke, Frank; Carmeliet, Peter; Simons, Michael  
CORPORATE SOURCE: Angiogenesis Research Center, Department of Surgery both at Beth Israel Deaconess Medical Center and Harvard Medical School, Boston, MA, 02215, USA  
SOURCE: Nature Medicine (New York) (2000), 6(1), 49-55  
CODEN: NAMEFI; ISSN: 1078-8956  
PUBLISHER: Nature America  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
AB Although tissue injury and inflammation are considered essential for the induction of **angiogenesis**, the mol. controls of this cascade are mostly unknown. Here we show that a macrophage-derived peptide,

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**PR39**, inhibited the ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein, resulting in accelerated formation of vascular structures in vitro and increased myocardial vasculature in mice. For the latter, coronary flow studies demonstrated that **PR39**-induced **angiogenesis** resulted in the prodn. of functional blood vessels. These findings show that **PR39** and related compds. can be used as potent inducers of **angiogenesis**, and that selective inhibition of hypoxia-inducible factor-1.alpha. degrdn. may underlie the mechanism of inflammation-induced **angiogenesis**.

IT 139637-11-9, **PR-39**

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(**PR39** peptide in regulation of **angiogenesis** by inhibiting ubiquitin-proteasome-dependent degrdn. of hypoxia-inducible factor-1.alpha. protein)

RN 139637-11-9 HCAPLUS

CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolylglycyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

REFERENCE COUNT: 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1996:357168 HCAPLUS

DOCUMENT NUMBER: 125:26311

TITLE: Synducin (syndecan expression-inducers) mediate modulation of tissue repair

INVENTOR(S): Gallo, Richard L.; Bernfield, Merton

PATENT ASSIGNEE(S): Children's Medical Center Corporation, USA

SOURCE: PCT Int. Appl., 32 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9609322	A2	19960328	WO 1995-US12080	19950922
WO 9609322	A3	19960523		
W: AU, CA, JP, KR				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5654273	A	19970805	US 1994-310722	19940922
AU 9538228	A1	19960409	AU 1995-38228	19950922
US 5863897	A	19990126	US 1996-728333	19961010
PRIORITY APPLN. INFO.:			US 1994-310722	19940922
			WO 1995-US12080	19950922

AB The membrane-permeating antibacterial peptide, **PR-39**, previously found only in the intestine, was purified from wound fluid and shown to possess syndecan-1 and syndecan-4 inductive activity specifically in mesenchymal cells. This is a newly recognized function that defines

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peptide contg. syndecan-inducing activity, and that are known as synducins. Therefore, a mol. with both antimicrobial and synducin activities is deposited in wounds where it can simultaneously reduce infection and the influence the action of growth factors, matrix components, and other cellular effectors involved in wound repair. Synducins, including **PR-39**, and derivs. thereof, as well as other proline and arginine-rich antimicrobial peptides, collectively referred to herein as "synducins", are therefore useful in the modulation of wound healing, as well as other disorders involving mesenchymal cells and cell surface mol. interaction, including metastatic disease, **angiogenesis**, restenosis, stasis or decubitis ulcers, and prevention of keloids.

IT 139637-11-9

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(synducins are syndecan expression-inducing peptides that mediate modulation of mesenchymal tissue repair)

RN 139637-11-9 HCAPLUS

CN L-Prolinamide, L-arginyl-L-arginyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-tyrosyl-L-leucyl-L-prolyl-L-arginyl-L-prolyl-L-arginyl-L-prolyl-L-prolyl-L-prolyl-L-phenylalanyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-leucyl-L-prolyl-L-prolyl-L-arginyl-L-isoleucyl-L-prolyl-L-prolyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl-L-prolyl-L-prolyl-L-arginyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

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GenCore version 5.1.4 p5 4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 13, 2003, 10:32:27 ; Search time 35 Seconds  
(without alignments)  
57.107 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPPPLPRPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	90	100.0	15	21	PR-39 derived angi
2	90	100.0	15	22	Amino acid sequenc
3	90	100.0	15	22	PR-39 derived pept
4	90	100.0	19	17	Leukocyte O2- prod
5	90	100.0	26	17	Leukocyte O2- prod
6	90	100.0	26	19	proline/Arginine r
7	90	100.0	39	14	Antibacterial pept
8	90	100.0	39	17	Leukocyte O2- prod
9	90	100.0	39	17	Synducin peptide (
10	90	100.0	39	17	Magainin-derived a

11	90	100.0	39	19	AAW75722	Proline/Arginine r
12	90	100.0	39	21	AAW75722	PR-39 peptide used
13	90	100.0	39	22	AAW75722	Amino acid sequenc
14	90	100.0	39	22	AAW75722	PR-39 peptide. Un
15	90	100.0	42	23	AAW75722	Antimicrobial pept
16	90	100.0	44	22	AAW75722	E. coli AMP gene p
17	83	92.2	14	17	AAW01450	Leukocyte O2- prod
18	83	92.2	14	19	AAW75725	proline/Arginine r
19	75	83.3	23	17	AAW01451	Leukocyte O2- prod
20	66	73.3	18	16	AAW75721	Bactenecin peptide
21	66	73.3	20	19	AAW75730	Proline/Arginine r
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25	66	73.3	59	21	AAW91699	Cationic peptide B
26	66	73.3	60	23	AAW07713	Antimicrobial pept
27	66	73.3	62	22	AAW51197	E. coli AMP gene B
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29	64	71.1	11	22	AAW26886	Amino acid sequenc
30	64	71.1	11	22	AAW26886	PR-39 derived pept
31	63	70.0	91	22	AAW61229	Propionibacterium
32	61	67.8	336	17	AAW05520	HCWV Toledo strain
33	60	66.7	953	23	AAU74761	Human protease PKT
34	60	66.7	953	23	AAU82708	Amino acid sequenc
35	59	65.6	39	21	AAW44779	Human secreted pro
36	59	65.6	59	17	AAW94448	Synducin peptide (
37	56.5	62.8	74	22	ABW42619	Peptide #10125 enc
38	56.5	62.8	74	22	AAW63510	Human brain expres
39	56.5	62.8	74	22	AAW76324	Human bone marrow
40	56.5	62.8	74	22	AAW36433	Peptide #10470 enc
41	56.5	62.8	74	23	ABG45621	Human peptide enco
42	56	62.2	692	22	AAU04851	Micromonospora eve
43	55.5	61.7	497	20	AAW04972	Mycobacterium spec
44	55	61.1	45	22	AAW18621	Peptide #5055 enco
45	54	60.0	87	22	AAU53075	Propionibacterium

ALIGNMENTS

RESULT 1  
AAW26885  
ID AAW26885 standard; peptide; 15 AA.  
XX AAW26885;  
XX AAW26885;  
DT 01-FEB-2001 (first entry)  
XX PR-39 derived angiogenesis regulatory peptide 1.  
DE PR-39 derived angiogenesis regulatory peptide 1.  
XX Angiogenesis; stimulation; PR-39; anoxia, myocardial infarction;  
KW myocardial ischaemia; proteasome.  
XX Synthetic.  
XX WO2000057895-A1.  
XX 05-OCT-2000.  
XX 16-MAR-2000; 2000WO-US07050.  
XX 26-MAR-1999; 99US-0276868.  
XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.  
XX Simons M, Gao Y;  
XX WPI; 2000-628319/60.  
XX Stimulating angiogenesis in situ, useful e.g. for treating anoxia and  
PT infarction, by administering a PR-39 oligopeptide that regulates  
PT enzymatic activity of proteasomes

PS Claim 12; Page 40; 51pp; English.

CC This invention relates to a method for the stimulation of angiogenesis in

CC situ within a targeted collection of viable cells. The method comprises

CC introducing, into the cytoplasm, at least 1 member of the PR-39

CC oligopeptide collective, which interacts with cytoplasmic proteasomes.

CC Part of the proteolytic activity of the proteasomes is selectively

CC altered so as to stimulate angiogenesis. The method is used to induce

CC angiogenesis in tissue that has suffered anoxia or infarction,

CC e.g. myocardial infarction or chronic myocardial ischaemia, and also to

CC study the mechanisms that control angiogenesis. The present sequence

CC represents a PR-39 derived peptide which interacts with the proteasome

CC and can be used in the method of the invention.

XX SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 21; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPP 15

Db 1 RRRPRPPYLP RPP 15

|||||

RESULT 2

AAB84691

ID AAB84591 standard; peptide; 15 AA.

XX AC AAB84691;

XX DT 17-SEP-2001 (first entry)

XX DE Amino acid sequence of a PR-39 derived peptide (residues 1-15).

XX KW PR-39; IkappaBalpha degradation; NFkappaB transcription factor;

XX KW myocardial infarction; chronic myocardial ischemia; heart disease;

XX KW anoxia.

XX OS Unidentified.

XX PN WO200147540-A1.

XX PD 05-JUL-2001.

XX PF 27-DEC-2000; 2000WO-US35293.

XX PR 29-DEC-1999; 99US-0474967.

XX PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX PI Simons M, Gao Y;

XX WPI; 2001-441690/47.

XX DR Selective inhibition of IkappaBalpha degradation within targeted viable

XX PT cell collection, involves interacting PR-39 oligopeptide with

XX PT IkappaBalpha and proteasomes, and altering proteolytic activity of

XX PT proteasomes.

XX PS Claim 11; Page 58; 69pp; English.

XX CC The present sequence represents a PR-39 derived peptide. It is used

XX CC for selective inhibition of IkappaBalpha degradation within a targeted

XX CC cell collection in-situ. The method is useful for selectively inhibiting

XX CC IkappaBalpha protein degradation in situ, decreasing the activity of

XX CC NFkappaB transcription factor and selective control of NFkappaB-dependent

XX CC gene expression in situ. The PR-39 derived peptides are useful in the

XX CC treatment of myocardial infarction, chronic myocardial ischemia of

XX CC heart disease and anoxia.

XX SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPP 15

Db 1 RRRPRPPYLP RPP 15

|||||

RESULT 3

AAB97277

ID AAB97277 standard; peptide; 15 AA.

XX AC AAB97277;

XX DT 09-AUG-2001 (first entry)

XX DE PR-39 derived peptide PR-15.

XX KW PR-39; cathelin; inflammation; wound healing; myocardial infarction;

XX KW proteasome; proteolysis; alpha7; peptide degradation; angiogenesis;

XX KW anoxia; chronic myocardial ischaemia; heart tissue.

XX OS Unidentified.

XX PN WO200130368-A1.

XX PD 03-MAY-2001.

XX PF 06-OCT-2000; 2000WO-US27552.

XX PR 25-OCT-1999; 99US-0426011.

XX PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.

XX PI Simons M, Gao Y;

XX WPI; 2001-355179/37.

XX DR Stimulation of angiogenesis and inhibition of proteasome mediated

XX PT degradation in cells, by introduction of PR-39 oligopeptide or its

XX PT N-terminal fragments or their conjugates, for use in anoxia and

XX PT infarction conditions.

XX PS Claim 12; Page 42; 52pp; English.

XX CC Peptides AAB97277 - AAB97279 represent PR-39 derived oligopeptides. PR-39

XX CC is a member of the the cathelin family of proteins, mature PR-39 is 39

XX CC amino acids in length (see AAB97280), and has been shown to play a role

XX CC in several inflammatory events including wound healing and myocardial

XX CC infarction. The PR-39 derived family of oligopeptides cause selective

XX CC inhibition of proteasome mediated degeneration of peptides and

XX CC stimulation of angiogenesis after their intracellular introduction to a

XX CC target cell. PR-39 derived peptides are able to interact with at least

XX CC the alpha7 subunit of the proteasomes, and therefore alter the

XX CC proteolytic activity of the proteasomes such that a selective increased

XX CC expression of specific proteins occurs. The invention includes methods

XX CC for the selective inhibition of proteasome mediated peptide degradation.

XX CC The method provides means for stimulating angiogenesis as required in

XX CC living tissues and organs which have suffered defects or have undergone

XX CC anoxia and/or infarction, myocardial infarction or chronic myocardial

XX CC ischaemia of heart tissue. Examples are the myocardium, skeletal or

XX CC smooth muscle, artery or vein, lung, brain, kidney, spleen, liver

XX CC gastrointestinal or nerve tissues, limbs, and extremities. A particular

XX CC example is after myocardial infarction or ischaemia.

XX SQ Sequence 15 AA;

Query Match 100.0%; Score 90; DB 22; Length 15;

Best Local Similarity 100.0%; Pred. No. 0.0002;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPP 15

Db 1 RRRPRPPLPRPRPP 15  
|||||

## RESULT 4

AAW01452  
ID AAW01452 standard; peptide; 19 AA.

AC AAW01452;

DT 18-JUN-1997 (first entry)

XX Leukocyte O2- production inhibitor peptide PR19.

XX Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;  
XX antimicrobial peptide; small intestine; human; neutrophil; bacteria;  
KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;  
KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;  
KW tissue damage; oxygen radical; inflammatory disease; therapy.

XX Synthetic.

XX WO9632129-A1.

XX 17-OCT-1996.

XX 10-APR-1996; 96WO-US04674.

XX 10-APR-1995; 95US-0419066.

XX (UNIV ) UNIV KANSAS STATE RES FOUND.

XX Blecha F, Shi J;

XX WPI; 1996-476842/47.

XX Inhibition of leukocyte super:oxide anion prodn. and attraction of  
PT leukocytes - using peptide(s) partic. based on antimicrobial PR-39

PS Disclosure; Page 27; 45pp; English.

XX AAW01447-W01454 represent fragments of the proline-arginine rich  
CC antimicrobial peptide PR39 (see AAW01446). The PR39 sequence was first  
CC isolated from porcine small intestine, and has also been identified in  
CC human and porcine neutrophils. PR39 kills bacteria by interfering with  
CC DNA and/or protein synthesis. PR39 also induces syndecan expression on  
CC mesenchymal cells. Syndecans are important in wound repair, showing that  
CC PR39 can be used in wound repair, as well as in antibacterial agents.  
CC These sequences, and PR39, can be used in the method of the invention.  
CC The method of the invention is for inhibiting leukocyte superoxide anion  
CC peptide (such as this sequence) capable of inhibiting leukocyte O2-  
CC production. The peptides can be used as medicaments for fighting  
CC infection by attracting leukocytes to a wound site and restricting  
CC tissue damage at the wound site caused by excessive oxygen radicals  
CC produced by these leukocytes. They can also be used to develop products  
CC for treating inflammatory disease states.

XX Sequence 19 AA;

Query Match 100.0%; Score 90; DB 17; Length 19;  
Best Local Similarity 100.0%; Pred. No. 0.00025;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPRPP 15  
|||||

Db 1 RRRPRPPLPRPRPP 15

## RESULT 5

AAW01447  
ID AAW01447 standard; peptide; 26 AA.

XX

AC AAW01447;

DT 18-JUN-1997 (first entry)

DE Leukocyte O2- production inhibitor peptide PR26.

XX Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;  
KW antimicrobial peptide; small intestine; human; neutrophil; bacteria;  
KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;  
KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;  
KW tissue damage; oxygen radical; inflammatory disease; therapy.

XX Synthetic.

XX WO9632129-A1.

XX 17-OCT-1996.

XX 10-APR-1996; 96WO-US04674.

XX 10-APR-1995; 95US-0419066.

XX (UNIV ) UNIV KANSAS STATE RES FOUND.

XX Blecha F, Shi J;

XX WPI; 1996-476842/47.

XX Inhibition of leukocyte super:oxide anion prodn. and attraction of  
PT leukocytes - using peptide(s) partic. based on antimicrobial PR-39

PS Claim 3; Page 26; 45pp; English.

XX AAW01447-W01454 represent fragments of the proline-arginine rich  
CC antimicrobial peptide PR39 (see AAW01446). The PR39 sequence was first  
CC isolated from porcine small intestine, and has also been identified in  
CC human and porcine neutrophils. PR39 kills bacteria by interfering with  
CC DNA and/or protein synthesis. PR39 also induces syndecan expression on  
CC mesenchymal cells. Syndecans are important in wound repair, showing that  
CC PR39 can be used in wound repair, as well as in antibacterial agents.  
CC These sequences, and PR39, can be used in the method of the invention.  
CC The method of the invention is for inhibiting leukocyte superoxide anion  
CC peptide (such as this sequence) capable of inhibiting leukocyte O2-  
CC production. The peptides can be used as medicaments for fighting  
CC infection by attracting leukocytes to a wound site and restricting  
CC tissue damage at the wound site caused by excessive oxygen radicals  
CC produced by these leukocytes. They can also be used to develop products  
CC for treating inflammatory disease states.

XX Sequence 26 AA;

Query Match 100.0%; Score 90; DB 17; Length 26;  
Best Local Similarity 100.0%; Pred. No. 0.00033;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPRPP 15  
|||||

Db 1 RRRPRPPLPRPRPP 15

## RESULT 6

AAW75723  
ID AAW75723 standard; peptide; 26 AA.

XX AAW75723;

DT 19-NOV-1998 (first entry)

DE Proline/Arginine rich peptide PR-26.

XX Proline; arginine; peptide; reperfusion injury; neutrophil; endothelium;  
KW superoxide; neutrophil superoxide anion; chemotaxis; NADPH oxidase;

KW coronary bypass; organ transplantation surgery.  
 XX Synthetic.  
 OS  
 XX  
 PN WO9835690-A1.  
 XX  
 XX  
 PD 20-AUG-1998.  
 XX  
 XX 17-FEB-1998; 98WO-US03207.  
 PF  
 XX 16-FEB-1998; 98US-0024975.  
 PR  
 PR 18-FEB-1997; 97US-0802306.  
 XX  
 XX (UNIV ) UNIV KANSAS STATE RES FOUND.  
 PA  
 XX  
 XX Blecha F, Ross CR, Shi J;  
 PI  
 XX WPI; 1998-495359/42.  
 XX  
 XX Reduction of reperfusion injury in temporarily occluded blood  
 PT vessels - by administration of a peptide which is rich in proline  
 PT or arginine residues  
 XX  
 PS Claim 3; Page 14-15; 35pp; English.  
 XX  
 CC Sequences AAW75722-W75732 are proline/arginine rich peptides that upon  
 CC administration into a mammal's bloodstream reduce reperfusion injury  
 CC (production of reactive oxygen species, neutrophil adherence to  
 CC endothelium, and extravasation of neutrophils). These peptides have two  
 CC requirements: they contain the consensus sequence PXXP, where P is a  
 CC proline residue and X is any amino acid residue, which has been found to  
 CC inhibit superoxide production, and secondly they have arginine residues  
 CC adjacent to these motifs, required for effective inhibition. It was  
 CC established by structural and function analysis that a peptide should  
 CC ideally contain 4 or 6 of these motifs, and that inhibitory activity is  
 CC correlated with the increase of length of peptides. The effectiveness  
 CC of these peptides was determined by investigating the production of the  
 CC neutrophil superoxide anion, and also the inhibition of neutrophil  
 CC chemotaxis. From this, it was found that all of the peptides inhibited  
 CC NADPH oxidase to some extent. All of the peptides also inhibit  
 CC neutrophil oxidase activity. PR-39 is believed, to be the most potent  
 CC endogenous down regulator of NADPH oxidase yet discovered, and from the  
 CC data produced, it can be suggested to be involved in eliminating or  
 CC reducing the reperfusion injury induced adhesion and extraction of  
 CC neutrophils. The peptides are also useful in connection with surgical  
 CC procedures such as coronary bypass and organ transplantation surgery.  
 XX  
 SQ Sequence 26 AA;  
 Query Match 100.0%; Score 90; DB 19; Length 26;  
 Best Local Similarity 100.0%; Pred. No. 0.00033;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 RRRPRPPYLP RPP 15  
 |||||  
 Db 1 RRRPRPPYLP RPP 15  
 |||||  
 RESULT 7  
 AAR30491  
 ID AAR30491 standard; peptide; 39 AA.  
 XX  
 AC AAR30491;  
 XX  
 XX 12-MAY-1993 (first entry)  
 DT  
 XX Antibacterial peptide.  
 DE  
 XX Pig; small intestine; endocrine; gram negative; bacteria; therapeutic;  
 KW veterinary medicine; prophylactic.  
 KW  
 XX Sus scrofa domestica.  
 OS  
 XX

PN WO9222578-A.  
 XX  
 PD 23-DEC-1992.  
 XX  
 XX 10-JUN-1992; 92WO-SE00394.  
 PF  
 XX 14-JUN-1991; 91SE-0001838.  
 PR  
 XX (BOMA/) BOMAN H G.  
 PA (JOER/) JOERNVALL H.  
 PA (LEEJ/) LEE J.  
 PA (MUTT/) MUTT V.  
 XX  
 PI Boman HG, Joernvall H, Lee J, Mutt V;  
 XX  
 XX WPI; 1993-018080/02.  
 DR  
 XX New anti-bacterial polypeptide - active against Gram negative  
 PT bacteria  
 PT  
 XX Claim 1; Page 10; 15pp; English.  
 PS  
 XX This peptide was isolated from the small intestine of a pig. The  
 CC small intestine is an important endocrine organ and many  
 CC physiologically active peptides have been isolated from it. This  
 CC peptide inhibits the growth of, and may kill, bacteria, pref. gram  
 CC negative bacteria. This peptide or its functional derivatives may be  
 CC used in human or veterinary medicine for therapeutic or prophylactic  
 CC use.  
 CC  
 XX Sequence 39 AA;  
 SQ  
 Query Match 100.0%; Score 90; DB 14; Length 39;  
 Best Local Similarity 100.0%; Pred. No. 0.00047;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 RRRPRPPYLP RPP 15  
 |||||  
 Db 1 RRRPRPPYLP RPP 15  
 |||||  
 RESULT 8  
 AAW01446  
 ID AAW01446 standard; peptide; 39 AA.  
 XX  
 AC AAW01446;  
 XX  
 XX 18-JUN-1997 (first entry)  
 DT  
 XX Leukocyte O2- production inhibitor peptide PR39.  
 DE  
 XX Inhibitor; leukocyte O2- production; proline-arginine rich peptide; pig;  
 KW antimicrobial peptide; small intestine; human; neutrophil; bacteria;  
 KW DNA synthesis; protein synthesis; inhibitor; syndecan expression;  
 KW mesenchymal cell; wound repair; superoxide anion; infection; leukocyte;  
 KW tissue damage; oxygen radical; inflammatory disease; therapy.  
 XX  
 OS Synthetic.  
 OS  
 XX WO9632129-A1.  
 PN  
 XX 17-OCT-1996.  
 PD  
 XX 10-APR-1996; 96WO-US04674.  
 PF  
 XX 10-APR-1995; 95US-0419066.  
 PR  
 XX (UNIV ) UNIV KANSAS STATE RES FOUND.  
 PA  
 XX Blecha F, Shi J;  
 PI  
 XX WPI; 1996-476842/47.  
 XX

PT Inhibition of leukocyte super:oxide anion prodn. and attraction of  
 XX leukocytes - using peptide(s) partic. based on antimicrobial PR-39  
 PS Claim 2; Page 26; 45pp; English.  
 XX This sequence represents the proline-arginine rich antimicrobial peptide  
 CC PR39. The PR39 sequence was first isolated from porcine small intestine,  
 CC and has also been identified in human and porcine neutrophils. PR39  
 CC kills bacteria by interfering with DNA and/or protein synthesis. PR39  
 CC also induces syndecan expression on mesenchymal cells. Syndecans are  
 CC important in wound repair, showing that PR39 can be used in wound repair,  
 CC as well as in antibacterial agents. This sequence, and the fragments of  
 CC it shown in AA00147-W01454, can be used in the method of the invention.  
 CC The method of the invention is for inhibiting leukocyte superoxide anion  
 CC (O2-) production. The method comprises administering to a leukocyte a  
 CC peptide (such as this sequence) capable of inhibiting leukocyte O2-  
 CC production. The peptides can be used as medicaments for fighting  
 CC infection by attracting leukocytes to a wound site and restricting tissue  
 CC damage at the wound site caused by excessive oxygen radicals produced by  
 CC these leukocytes. They can also be used to develop products for treating  
 CC inflammatory disease states.  
 XX  
 SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 17; Length 39;  
 Best Local Similarity 100.0%; Pred. No. 0.00047;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 RRRPRPPYLP RPP 15  
 |||||  
 DB 1 RRRPRPPYLP RPP 15

RESULT 9  
 AAR94446  
 ID AAR94446 standard; peptide; 39 AA.  
 AC AAR94446;  
 XX  
 XX 05-NOV-1996 (first entry)  
 DT  
 XX Syndecan peptide (PR-39) induces syndecan expression.  
 DE  
 XX Syndecan; induction; expression; syndecan-1; syndecan-4; surface;  
 KW mesenchymal cell; fibroblast; epithelial; PR-39; treatment; stasis;  
 KW decubitus; ulcers; keloids; skin burns; ischemic tissues;  
 KW hypercoagulation states; prevention; tumour metastasis; restenosis;  
 KW inhibition; angiogenesis; proliferation; endothelial.  
 XX  
 OS Synthetic.  
 XX WO9609322-A2.  
 PN  
 XX 28-MAR-1996.  
 PD  
 XX 22-SEP-1995; 95WO-US12080.  
 PF  
 XX 22-SEP-1994; 94US-0310722.  
 PR  
 XX (CHIL-) CHILDRENS MEDICAL CENT.  
 PA  
 XX Bernfield M, Gallo RL;  
 PI  
 XX WPI; 1996-188401/19.  
 DR  
 XX Modulating mesenchymal interaction by administration of syndecan -  
 PT used in the treatment of wounds, tumours, restenosis, etc  
 XX  
 PS Claim 4; Page 26; 34pp; English.  
 XX The present peptide is a syndecan, which induces the expression of  
 CC syndecan-1 and syndecan-4 on the surface of mesenchymal cells, esp.  
 CC fibroblasts and epithelial cells. The 36 N-terminal amino acids of

CC the peptide were found to be identical to the 36 N-terminal amino  
 CC acids of PR-39, a Pro and Arg rich antibacterial peptide previously  
 CC found in porcine intestine (WO9222578). Syndecins may be used in  
 CC the treatment of stasis and decubitus ulcers, keloids, skin burns,  
 CC ischemic tissues and hypercoagulation states, prevention of tumour  
 CC metastasis, restenosis inhibition and endothelial cell angiogenesis  
 CC and proliferation induction.  
 CC Human microvascular endothelial cells were assayed for syndecan-4  
 CC expression following exposure to 5 % wound fluid, dbcAMP (1 mM),  
 CC the present peptide (10 microm) or a blank, to give respective  
 CC cell surface syndecan-4 values (MOD/m in) of approx. 1.75, 1.70,  
 CC 1.80 and 0.95.  
 XX

SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 17; Length 39;

Best Local Similarity 100.0%; Pred. No. 0.00047;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 RRRPRPPYLP RPP 15  
 |||||  
 DB 1 RRRPRPPYLP RPP 15

RESULT 10  
 AAR99121  
 ID AAR99121 standard; peptide; 39 AA.  
 XX  
 AC AAR99121;  
 XX  
 XX 28-OCT-1996 (first entry)  
 DT  
 XX Magainin-derived antimicrobial STD-inhibiting peptide, MSI-1312.

DE  
 XX STD; sexually transmitted disease; HIV; human immunodeficiency virus;  
 KW herpes simplex virus; HSV; Neisseria gonorrhoeae; Candida; Chlamydia;  
 KW magainin; antimicrobial; squalamine.  
 XX  
 OS Synthetic.  
 XX  
 XX Key Location/Qualifiers  
 FH Modified-site 39  
 FT /note= "amidated"  
 FT  
 XX WO9608270-A2.  
 PN  
 XX 21-MAR-1996.  
 PD  
 XX 13-SEP-1995; 95WO-US11675.  
 PF  
 XX 13-SEP-1994; 94US-0305475.  
 PR  
 XX (MAGA-) MAGAININ PHARM INC.  
 PA  
 XX Bedi G, Jacob L, Williams T, Zasloff M;  
 PI  
 XX WPI; 1996-179725/18.  
 DR  
 XX Inhibiting sexually transmitted disease e.g. HIV or herpes simplex -  
 PT by administering magainin antimicrobial or squalamine cpd. to  
 PT inhibit transmission  
 XX  
 PS Example 1; Page 32; 60pp; English.  
 XX  
 XX AAR99116-R99123 are antimicrobial, magainin-analogue peptides that may  
 CC be used to treat sexually transmitted diseases (STDs) caused by  
 CC Chlamydia, HIV, herpes simplex virus, Neisseria gonorrhoeae or  
 CC Candida infection. The peptides inhibit STDs by either killing the  
 CC infectious organism, impeding the infection mechanism or  
 CC interrupting the replication cycle of the organism. Squalamine (an  
 CC aminosterol host defence molecule of the dog fish shark Squalus  
 CC acanthias) and PGla (a frog antimicrobial peptide) analogues may  
 CC also be useful in inhibiting STD infection and transmission.

```

XX SQ Sequence 39 AA;
Query Match 100.0%; Score 90; DB 17; Length 39;
Best Local Similarity 100.0%; Pred. No. 0.00047;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
Db 1 RRRPRPPYLPRPRPP 15

RESULT 11
AAW75722
ID AAW75722 standard; peptide; 39 AA.
XX
AC AAW75722;
XX
DT 19-NOV-1998 (first entry)
XX
DE Proline/Arginine rich peptide PR-39.
XX
KW Proline; arginine; peptide; reperfusion injury; neutrophil; endothelium;
KW superoxide; neutrophil superoxide anion; chemotaxis; NADPH oxidase;
KW coronary bypass; organ transplantation surgery.
XX
OS Synthetic.
XX
PN WO9835690-A1.
XX
PD 20-AUG-1998.
XX
PF 17-FEB-1998; 98NO-US03207.
XX
PR 16-FEB-1998; 98US-0024975.
PR 18-FEB-1997; 97US-0802306.
XX
PA (UNIV ) UNIV KANSAS STATE RES FOUND.
XX
PI Blecha F, Ross CR, Shi J;
XX
DR WPI; 1998-495359/42.
XX
PT Reduction of reperfusion injury in temporarily occluded blood
PT vessels - by administration of a peptide which is rich in proline
PT or arginine residues
XX
Claim 3; Page 14; 35pp; English.
XX
CC Sequences AAW75722-W75732 are proline/arginine rich peptides that upon
CC administration into a mammal's bloodstream reduce reperfusion injury
CC (production of reactive oxygen species, neutrophil adherence to
CC endothelium, and extravasation of neutrophils). These peptides have two
CC requirements: they contain the consensus sequence PXXP, where P is a
CC proline residue and X is any amino acid residue, which has been found to
CC inhibit superoxide production, and secondly they have arginine residues
CC adjacent to these motifs, required for effective inhibition. It was
CC established by structural and function analysis that a peptide should
CC ideally contain 4 or 6 of these motifs, and that inhibitory activity is
CC correlated with the increase of length of peptides. The effectiveness
CC of these peptides was determined by investigating the production of the
CC neutrophil superoxide anion, and also the inhibition of neutrophil
CC chemotaxis. From this, it was found that all of the peptides inhibited
CC NADPH oxidase to some extent. All of the peptides also inhibit
CC neutrophil oxidase activity. PR-39 is believed, to be the most potent
CC endogenous down regulator of NADPH oxidase yet discovered, and from the
CC data produced, it can be suggested to be involved in eliminating or
CC reducing the reperfusion injury induced adhesion and extraction of
CC neutrophils. The peptides are also useful in connection with surgical
CC procedures such as coronary bypass and organ transplantation surgery.
XX
SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 19; Length 39;
Best Local Similarity 100.0%; Pred. No. 0.00047;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
Db 1 RRRPRPPYLPRPRPP 15

RESULT 12
AAB26888
ID AAB26888 standard; peptide; 39 AA.
XX
AC AAB26888;
XX
DT 01-FEB-2001 (first entry)
XX
DE PR-39 peptide used in angiogenesis control.
XX
KW Angiogenesis; stimulation; PR-39; anoxia; myocardial infarction;
KW myocardial ischaemia; proteasome.
XX
OS Synthetic.
XX
PN WO200057895-A1.
XX
PD 05-OCT-2000.
XX
PF 16-MAR-2000; 2000WO-US07050.
XX
PR 26-MAR-1999; 99US-0276868.
XX
PA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
XX
PI Simons M, Gao Y;
XX
DR WPI; 2000-628319/60.
XX
PT Stimulating angiogenesis in situ, useful e.g. for treating anoxia and
PT infarction, by administering a PR-39 oligopeptide that regulates
PT enzymatic activity of proteasomes -
XX
PS Disclosure; Page 21; 51pp; English.
XX
CC This invention relates to a method for the stimulation of angiogenesis in
CC situ within a targeted collection of viable cells. The method comprises
CC introducing, into the cytoplasm, at least 1 member of the PR-39
CC oligopeptide collective, which interacts with cytoplasmic proteasomes.
CC Part of the proteolytic activity of the proteasomes is selectively
CC altered so as to stimulate angiogenesis. The method is used to induce
CC angiogenesis in tissue that has suffered anoxia or infarction,
CC e.g. myocardial infarction or chronic myocardial ischaemia, and also to
CC study the mechanisms that control angiogenesis. The present sequence
CC represents the PR-39 peptide from which peptide used in the method of
CC the invention are derived.
XX
SQ Sequence 39 AA;

Query Match 100.0%; Score 90; DB 21; Length 39;
Best Local Similarity 100.0%; Pred. No. 0.00047;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLPRPRPP 15
Db 1 RRRPRPPYLPRPRPP 15

RESULT 13
AAB84690
ID AAB84690 standard; protein; 39 AA.
XX
AC AAB84690;
XX

```



DT 17-SEP-2001 (first entry)  
 XX Amino acid sequence of a PR-39 protein.  
 DE PR-39; IkappaBalpa degradation; NFkappaB transcription factor;  
 KW myocardial infarction; chronic myocardial ischemia; heart disease;  
 KW anoxia.  
 XX Unidentified.  
 OS WO200147540-A1.  
 PN 05-JUL-2001.  
 XX 27-DEC-2000; 2000WO-US35293.  
 PF 29-DEC-1999; 99US-0474967.  
 XX (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.  
 PA Simons M, Gao Y;  
 PI WPI; 2001-441690/47.  
 DR Selective inhibition of IkappaBalpa degradation within targeted viable  
 XX cell collection, involves interacting PR-39 oligopeptide with  
 PT IkappaBalpa and proteasomes, and altering proteolytic activity of  
 PT proteasomes.  
 XX Disclosure; Page 30; 69pp; English.  
 PS The present sequence represents a PR-39 protein. The specification  
 CC describes PR-39 derived peptides, which are used for selective  
 CC inhibition of IkappaBalpa degradation within a targeted cell collection  
 CC in-situ. The method is useful for selectively inhibiting IkappaBalpa  
 CC protein degradation in situ, decreasing the activity of NFkappaB  
 CC transcription factor and selective control of NFkappaB-dependent gene  
 CC expression in situ. The PR-39 derived peptides are useful in the  
 CC treatment of myocardial infarction, chronic myocardial ischemia of  
 CC heart disease and anoxia.  
 XX Sequence 39 AA;  
 SQ Query Match 100.0%; Score 90; DB 22; Length 39;  
 Best Local Similarity 100.0%; Pred. No. 0.00047;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 RRRPRPPYLP RPP 15  
 DB 1 RRRPRPPYLP RPP 15  
 RESULT 14  
 AAB97280  
 ID AAB97280 standard; peptide; 39 AA.  
 AC AAB97280;  
 XX 09-AUG-2001 (first entry)  
 DT PR-39 peptide.  
 DE PR-39; cathelin; inflammation; wound healing; myocardial infarction;  
 KW proteasome; proteolysis; alpha7; peptide degradation; angiogenesis;  
 KW anoxia; chronic myocardial ischemia; heart tissue.  
 XX Unidentified.  
 OS WO200130368-A1.  
 PN 03-MAY-2001.  
 XX 06-OCT-2000; 2000WO-US27552.  
 PF

XX 25-OCT-1999; 99US-0426011.  
 PR (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.  
 PA Simons M, Gao Y;  
 PI WPI; 2001-355179/37.  
 DR Stimulation of angiogenesis and inhibition of proteasome mediated  
 XX degradation in cells, by introduction of PR-39 oligopeptide or its  
 PT N-terminal fragments or their conjugates, for use in anoxia and  
 PT infarction conditions.  
 XX Disclosure; Page 21; 52pp; English.  
 PS Peptides AAB97277 - AAB97279 represent PR-39 derived oligopeptides. PR-39  
 CC is a member of the the cathelin family of proteins, mature PR-39  
 CC represented by the present sequence is 39 amino acids in length, and has  
 CC been shown to play a role in several inflammatory events including wound  
 CC healing and myocardial infarction. The PR-39 derived family of  
 CC oligopeptides cause selective inhibition of proteasome mediated  
 CC degeneration of peptides and stimulation of angiogenesis after their  
 CC intracellular introduction to a target cell. PR-39 derived peptides are  
 CC able to interact with at least the alpha7 subunit of the proteasomes, and  
 CC therefore alter the proteolytic activity of proteasomes such that a  
 CC selective increased expression of specific proteins occurs. The invention  
 CC includes methods for the selective inhibition of proteasome mediated  
 CC peptide degradation. The method provides means for stimulating  
 CC angiogenesis as required in living tissues and organs which have suffered  
 CC defects or have undergone anoxia and/or infarction, myocardial infarction  
 CC or chronic myocardial ischemia of heart tissue. Examples are the  
 CC myocardium, skeletal or smooth muscle, artery or vein, lung, brain,  
 CC kidney, spleen, liver, gastrointestinal or nerve tissues, limbs, and  
 CC extremities. A particular example is after myocardial infarction or  
 CC ischaemia.  
 XX Sequence 39 AA;  
 SQ Query Match 100.0%; Score 90; DB 22; Length 39;  
 Best Local Similarity 100.0%; Pred. No. 0.00047;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 RRRPRPPYLP RPP 15  
 DB 1 RRRPRPPYLP RPP 15  
 RESULT 15  
 ABB07714  
 ID ABB07714 standard; peptide; 42 AA.  
 XX ABB07714;  
 AC 10-JUN-2002 (first entry)  
 DT Antimicrobial peptide PR-39 C-terminal fragment.  
 DE Vaccine; cathelicidin; antimicrobial; immunostimulant; immune response;  
 XX antigen presenting cell; adjuvant; porcine; PR-39.  
 KW Sus sp.  
 OS WO200213857-A2.  
 PN 21-FEB-2002.  
 PD 17-AUG-2001; 2001WO-EP09529.  
 PF 17-AUG-2000; 2000AT-0001416.  
 XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.  
 PA

PI Fritz J, Mattner F, Zauner W, Buschle M, Egyed A;  
 XX  
 DR WPI; 2002-269154/31.  
 XX  
 PT Vaccine for active immunization or for preparing an adjuvant for  
 PT enhancing an immune response to at least one antigen, comprises at  
 PT least one antigen and at least one cathelicidin derived antimicrobial  
 PT peptide -  
 XX  
 PS Disclosure; Fig 3; 65pp; English.  
 XX  
 CC The invention relates to a vaccine comprising at least one antigen and at  
 CC least one cathelicidin derived antimicrobial peptide or its derivative.  
 CC The vaccine is useful for active immunization, especially of humans or  
 CC animals without protection against the specific antigen. The cathelicidin  
 CC derived antimicrobial peptide is useful in the preparation of an adjuvant  
 CC for enhancing the immune response to at least one antigen, where the  
 CC adjuvant enhances the uptake of at least one antigen in antigen  
 CC presenting cells (APC), and the adjuvant is added to the vaccine.  
 CC Sequences ABB07708-15 represent C-terminal fragments of antimicrobial  
 CC peptides of the cathelicidin family.  
 XX  
 SQ Sequence 42 AA;  
 Query Match 100.0%; Score 90; DB 23; Length 42;  
 Best Local Similarity 100.0%; Pred. No. 0.0005;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 RRRPRPPYLP RPP 15  
 Db 1 RRRPRPPYLP RPP 15  
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Search completed: May 13, 2003, 10:40:32  
 Job time : 36 secs

09/426,011

GenCore version 5.1.4 p5 4578  
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:40:17 ; Search time 14 Seconds  
(without alignments)  
31.525 Million cell updates/sec

Title: US-09-426-011D-3

Perfect score: 90

Sequence: 1 RRRPPYLPRLPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 262574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents AA:\*

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2: /cgn2\_6/prodata/1/iaa/5B\_COMB.pep:\*

3: /cgn2\_6/prodata/1/iaa/6A\_COMB.pep:\*

4: /cgn2\_6/prodata/1/iaa/6B\_COMB.pep:\*

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6: /cgn2\_6/prodata/1/iaa/backfiles1.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	90	100.0	26	US-08-419-066-2	Sequence 2, Appli
2	90	100.0	26	US-09-024-975-2	Sequence 2, Appli
3	90	100.0	39	US-08-162-052-1	Sequence 1, Appli
4	90	100.0	39	US-08-310-722-1	Sequence 1, Appli
5	90	100.0	39	US-08-419-066-1	Sequence 1, Appli
6	90	100.0	39	US-08-728-333-1	Sequence 1, Appli
7	90	100.0	39	US-09-024-975-1	Sequence 1, Appli
8	90	100.0	39	PCT-US95-12080-1	Sequence 1, Appli
9	83	92.2	14	US-09-024-975-4	Sequence 4, Appli
10	66	73.3	20	US-09-024-975-9	Sequence 9, Appli
11	61	67.8	336	US-08-414-926A-26	Sequence 26, Appli
12	61	67.8	336	US-08-926-922-26	Sequence 26, Appli
13	61	67.8	336	US-09-253-682-26	Sequence 26, Appli
14	61	67.8	336	US-09-527-657-26	Sequence 26, Appli
15	59	65.6	59	PCT-US95-12080-3	Sequence 3, Appli
16	53	58.9	18	US-08-205-938A-23	Sequence 23, Appli
17	53	58.9	18	US-08-205-938A-24	Sequence 24, Appli
18	53	58.9	18	US-09-230-180-20	Sequence 20, Appli
19	53	58.9	18	PCT-US95-02626-23	Sequence 23, Appli
20	53	58.9	18	PCT-US95-02626-23	Sequence 23, Appli
21	52	57.8	18	US-08-205-938A-25	Sequence 25, Appli
22	52	57.8	18	PCT-US95-02626-25	Sequence 25, Appli
23	51.5	57.2	355	US-08-483-533-41	Sequence 41, Appli
24	51.5	57.2	355	US-09-283-471A-41	Sequence 41, Appli
25	51.5	57.2	355	PCT-US91-06532-3	Sequence 3, Appli
26	51	56.7	16	US-08-205-938A-8	Sequence 8, Appli
27	51	56.7	16	PCT-US95-02626-8	Sequence 8, Appli

28	51	56.7	180	3	US-09-187-331-5	Sequence 5, Appli
29	51	56.7	180	4	US-09-470-946-5	Sequence 5, Appli
30	51	56.7	195	3	US-09-187-331-1	Sequence 1, Appli
31	51	56.7	195	4	US-09-470-946-1	Sequence 1, Appli
32	50.5	56.1	169	4	US-08-483-533-28	Sequence 28, Appli
33	50.5	56.1	169	4	US-09-283-471A-28	Sequence 28, Appli
34	50.5	56.1	393	4	US-09-432-470-2	Sequence 2, Appli
35	50.5	56.1	393	4	US-09-432-470-4	Sequence 4, Appli
36	50	55.6	16	1	US-08-205-938A-7	Sequence 7, Appli
37	50	55.6	16	1	US-08-205-938A-28	Sequence 28, Appli
38	50	55.6	16	5	PCT-US95-02626-7	Sequence 7, Appli
39	50	55.6	16	5	PCT-US95-02626-28	Sequence 28, Appli
40	50	55.6	17	1	US-08-205-938A-27	Sequence 27, Appli
41	50	55.6	17	5	PCT-US95-02626-27	Sequence 27, Appli
42	49.5	55.0	129	4	US-09-199-637A-97	Sequence 97, Appli
43	49	54.4	26	4	US-09-024-975-8	Sequence 8, Appli
44	48.5	53.9	716	4	US-09-186-276B-67	Sequence 67, Appli
45	48.5	53.9	716	4	US-08-842-445-67	Sequence 67, Appli

ALIGNMENTS

RESULT 1

US-08-419-066-2  
; Sequence 2, Application US/08419066  
; Patent No, 5830993  
; GENERAL INFORMATION:  
; APPLICANT: Blecha, Frank  
; APPLICANT: Shi, Jishu  
; TITLE OF INVENTION: SYNTHETIC ANTIMICROBIAL PEPTIDE  
; NUMBER OF SEQUENCES: 2  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: John M. Collins, Hovey, Williams, Timmons &  
; ADDRESSEE: Collins  
; STREET: 2405 Grand Boulevard, Suite 400  
; CITY: Kansas City  
; STATE: Missouri  
; COUNTRY: U.S.A.  
; ZIP: 64108  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/419,066  
; FILING DATE:  
; CLASSIFICATION: 530  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Collins, John M.  
; REGISTRATION NUMBER: 26262  
; REFERENCE/DOCKET NUMBER: 23625  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (816) 474-9050  
; TELEFAX: (816) 474-9057  
; INFORMATION FOR SEQ ID NO: 2:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 26 amino acids  
; TYPE: amino acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: peptide  
; HYPOTHETICAL: NO  
; ANTI-SENSE: NO  
; FRAGMENT TYPE: N-terminal  
US-08-419-066-2

Query Match 100.0%; Score 90; DB 2; Length 26;  
Best Local Similarity 100.0%; Pred. No. 6.2e-05;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPPYLPRLPP 15



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TELEFAX: (404)-815-6555
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
PUBLICATION INFORMATION:
AUTHORS: Lee, Jong-Youn
AUTHORS: Boman, Hans G.
AUTHORS: Mutt, Viktor
AUTHORS: Jornvall, Hans
TITLE: No. 5654273el Polypeptides And Their Use
JOURNAL: PCT WO 92/22578
DATE: 12/23/92
RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
US-08-310-722-1

Query Match 100.0%; Score 90; DB 1; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPRPP 15
   |||||
Db 1 RRRPRPPLPRPRPP 15

RESULT 5
US-08-419-066-1
; Sequence 1, Application US/08419066
; Patent No. 5830993
; GENERAL INFORMATION:
; APPLICANT: Blecha, Frank
; APPLICANT: Shi, Jishu
; TITLE OF INVENTION: SYNTHETIC ANTIMICROBIAL PEPTIDE
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: John M. Collins, Hovey, Williams, Timmons &
; ADDRESSEE: Collins
; STREET: 2405 Grand Boulevard, Suite 400
; CITY: Kansas City
; STATE: Missouri
; COUNTRY: U.S.A.
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/419,066
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Collins, John M.
; REGISTRATION NUMBER: 26262
; REFERENCE/DOCKET NUMBER: 23625
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (816) 474-9050
; TELEFAX: (816) 474-9057
; INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
FRAGMENT TYPE: N-terminal
US-08-419-066-1
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TELEFAX: (404)-815-6555
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
PUBLICATION INFORMATION:
AUTHORS: Lee, Jong-Youn
AUTHORS: Boman, Hans G.
AUTHORS: Mutt, Viktor
AUTHORS: Jornvall, Hans
TITLE: No. 5863897el Polypeptides And Their Use
JOURNAL: PCT WO 92/22578
DATE: 12/23/92
RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
US-08-728-333-1

Query Match 100.0%; Score 90; DB 2; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPRPP 15
   |||||
Db 1 RRRPRPPLPRPRPP 15

RESULT 6
US-08-728-333-1
; Sequence 1, Application US/08728333
; Patent No. 5863897
; GENERAL INFORMATION:
; APPLICANT: Gallo, Richard L.
; APPLICANT: Klagsbrun, Michael
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabst
; STREET: 1100 Peachtree Street, Suite 2800
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-4530
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/728,333
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/310,722
; FILING DATE: 22-SEP-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Pabst, Patrea L.
; REGISTRATION NUMBER: 31,284
; REFERENCE/DOCKET NUMBER: CMCC379
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-815-6508
; TELEFAX: (404)-815-6555
; INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 39 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
PUBLICATION INFORMATION:
AUTHORS: Lee, Jong-Youn
AUTHORS: Boman, Hans G.
AUTHORS: Mutt, Viktor
AUTHORS: Jornvall, Hans
TITLE: No. 5863897el Polypeptides And Their Use
JOURNAL: PCT WO 92/22578
DATE: 12/23/92
RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
US-08-728-333-1

Query Match 100.0%; Score 90; DB 2; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPLPRPRPP 15
   |||||
Db 1 RRRPRPPLPRPRPP 15
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RESULT 7
US-09-024-975-1
; Sequence 1, Application US/09024975
; Patent No. 6133233
; GENERAL INFORMATION:
; APPLICANT: ROSS, CHRISTOPHER R.
; APPLICANT: BLECHA, FRANK
; APPLICANT: SHI, JISHU
; TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
; STREET: 2405 GRAND BLVD., SUITE 400
; CITY: KANSAS CITY
; STATE: MO
; COUNTRY: USA
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/024,975
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/802,306
; FILING DATE: 18-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: COLLINS, JOHN M.
; REGISTRATION NUMBER: 26,262
; REFERENCE/DOCKET NUMBER: 25585-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 816/474-9050
; TELEFAX: 816/474-9057
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-09-024-975-1

Query Match 100.0%; Score 90; DB 4; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRPP 15
| | | | | | | | | | | | | | |
DB 1 RRRPRPPYLP RPRPP 15

RESULT 8
PCT-US95-12080-1
; Sequence 1, Application PC/TUS9512080
; GENERAL INFORMATION:
; APPLICANT: Children's Medical Center Corporation
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Patrea L. Pabst
; STREET: 2800 One Atlantic Center
; STREET: 1201 West Peachtree
; CITY: Atlanta
; STATE: Georgia
; COUNTRY: USA
; ZIP: 30309-3450
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
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; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/12080
; FILING DATE:
; CLASSIFICATION:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (404)-873-8794
; TELEFAX: (404)-815-8795
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 39 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PUBLICATION INFORMATION:
; AUTHORS: Lee, Jong-Youn
; AUTHORS: Boman, Hans G.
; AUTHORS: Mutt, Viktor
; AUTHORS: Jornvall, Hans
; TITLE: Novel Polypeptides And Their Use
; JOURNAL: PCT WO 92/22578
; DATE: 12/23/92
; RELEVANT RESIDUES IN SEQ ID NO: 1: FROM 1 TO 39
; PCT-US95-12080-1

Query Match 100.0%; Score 90; DB 5; Length 39;
Best Local Similarity 100.0%; Pred. No. 9.1e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPRPP 15
| | | | | | | | | | | | | | |
DB 1 RRRPRPPYLP RPRPP 15

RESULT 9
US-09-024-975-4
; Sequence 4, Application US/09024975
; Patent No. 6133233
; GENERAL INFORMATION:
; APPLICANT: ROSS, CHRISTOPHER R.
; APPLICANT: BLECHA, FRANK
; APPLICANT: SHI, JISHU
; TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS
; STREET: 2405 GRAND BLVD., SUITE 400
; CITY: KANSAS CITY
; STATE: MO
; COUNTRY: USA
; ZIP: 64108
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/024,975
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/802,306
; FILING DATE: 18-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: COLLINS, JOHN M.
; REGISTRATION NUMBER: 26,262
; REFERENCE/DOCKET NUMBER: 25585-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 816/474-9050
; TELEFAX: 816/474-9057
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
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LENGTH: 14 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
US-09-024-975-4

Query Match 92.2%; Score 83; DB 4; Length 14;  
Best Local Similarity 100.0%; Pred. No. 0.00023;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPPPLPRPP 14  
DB 1 RRRPPPLPRPP 14

## RESULT 10

US-09-024-975-9  
Sequence 9, Application US/09024975  
Patent No. 6133233  
GENERAL INFORMATION:  
APPLICANT: ROSS, CHRISTOPHER R.  
APPLICANT: BLECHA, FRANK  
TITLE OF INVENTION: PEPTIDE MODULATION OF REPERFUSION INJURY  
NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HOVEY, WILLIAMS, TIMMONS & COLLINS  
STREET: 2405 GRAND BLVD., SUITE 400  
CITY: KANSAS CITY  
STATE: MO  
COUNTRY: USA  
ZIP: 64108  
COMPUTER READABLE FORM:  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/024,975  
FILING DATE:

## CLASSIFICATION:

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/802,306  
FILING DATE: 18-FEB-1997  
ATTORNEY/AGENT INFORMATION:  
NAME: COLLINS, JOHN M.  
REGISTRATION NUMBER: 26,262  
REFERENCE/DOCKET NUMBER: 25585-A  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 816/474-9050  
TELEFAX: 816/474-9057  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 20 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: peptide  
US-09-024-975-9

Query Match 73.3%; Score 66; DB 4; Length 20;  
Best Local Similarity 85.7%; Pred. No. 0.032;  
Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPPPLPRPP 14  
DB 2 RRRPPPLPRPP 15

## RESULT 11

US-08-414-926A-26  
Sequence 26, Application US/08414926A  
Patent No. 5721354  
GENERAL INFORMATION:

APPLICANT: Spaete, Richard  
APPLICANT: Cha, Tai-An  
TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS  
NUMBER OF SEQUENCES: 27  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum  
STREET: 5 Palo Alto Square  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94306-2155  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/414,926A  
FILING DATE: March 31, 1995  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Cserr, Luann  
REGISTRATION NUMBER: 31,822  
REFERENCE/DOCKET NUMBER: AVIR-011/OOUS  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-494-7622  
TELEFAX: 415-857-0663  
INFORMATION FOR SEQ ID NO: 26:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 336 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
IMMEDIATE SOURCE:  
CLONE: tol.22  
FEATURE:  
NAME/KEY: Protein  
LOCATION: 1..336  
OTHER INFORMATION: /label= UL151  
US-08-414-926A-26

Query Match 67.8%; Score 61; DB 1; Length 336;  
Best Local Similarity 78.6%; Pred. No. 1.7;  
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 RRRPPPLPRPP 15  
DB 279 RRRPPPLPRPP 292

## RESULT 12

US-08-926-922-26  
Sequence 26, Application US/08926922  
Patent No. 5925751  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard  
APPLICANT: Cha, Tai-An  
TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS  
NUMBER OF SEQUENCES: 27  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Luann Cserr Attorney at Law  
STREET: 750 Arimo Avenue  
CITY: Oakland  
STATE: CA  
COUNTRY: USA  
ZIP: 94610  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/926,922

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; FILING DATE: September 10, 1997
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Cserr, Luann
; REGISTRATION NUMBER: 31,822
; REFERENCE/DOCKET NUMBER: AVIR 11A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-834-1448
; TELEFAX: 510-839-7810
; INFORMATION FOR SEQ ID NO: 26:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 336 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; IMMEDIATE SOURCE:
; CLONE: tol.22
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..336
; OTHER INFORMATION: /label= UL151
; US-08-926-922-26
;
; Query Match 67.8%; Score 61; DB 2; Length 336;
; Best Local Similarity 78.6%; Pred. No. 1.7;
; Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
;
; QY 2 RRRPPYLP RPP 15
; Db 279 RRPPIQLRPP 292
;
; RESULT 13
; US-09-253-682-26
; Sequence 26, Application US/09253682
; Patent No. 6040170
; GENERAL INFORMATION:
; APPLICANT: Spaete, Richard
; APPLICANT: Cha, Tai-An
; TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Luann Cserr Attorney at Law
; STREET: 750 Arimo Avenue
; CITY: Oakland
; STATE: CA
; COUNTRY: USA
; ZIP: 94610
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/253,682
; FILING DATE: September 10, 1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/926,922
; FILING DATE: September 10, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Cserr, Luann
; REGISTRATION NUMBER: 31,822
; REFERENCE/DOCKET NUMBER: AVIR 11A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-834-1448
; TELEFAX: 510-839-7810
; INFORMATION FOR SEQ ID NO: 26:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 336 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; IMMEDIATE SOURCE:
; CLONE: tol.22
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..336
; OTHER INFORMATION: /label= UL151
; US-09-527-657-26
;
; Query Match 67.8%; Score 61; DB 4; Length 336;
; Best Local Similarity 78.6%; Pred. No. 1.7;
; Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
;
; QY 2 RRRPPYLP RPP 15
; Db 279 RRPPIQLRPP 292
;
; RESULT 14
; US-09-527-657-26
; Sequence 26, Application US/09527657
; Patent No. 6291236
; GENERAL INFORMATION:
; APPLICANT: Spaete, Richard
; APPLICANT: Cha, Tai-An
; TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Luann Cserr Attorney at Law
; STREET: 750 Arimo Avenue
; CITY: Oakland
; STATE: CA
; COUNTRY: USA
; ZIP: 94610
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/527,657
; FILING DATE: 17-Mar-2000
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/926,922
; FILING DATE: September 10, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Cserr, Luann
; REGISTRATION NUMBER: 31,822
; REFERENCE/DOCKET NUMBER: AVIR 11A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-834-1448
; TELEFAX: 510-839-7810
; INFORMATION FOR SEQ ID NO: 26:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 336 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; IMMEDIATE SOURCE:
; CLONE: tol.22
; FEATURE:
; NAME/KEY: Protein
; LOCATION: 1..336
; OTHER INFORMATION: /label= UL151
; US-09-527-657-26
;
; Query Match 67.8%; Score 61; DB 4; Length 336;
; Best Local Similarity 78.6%; Pred. No. 1.7;
; Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
;
; QY 2 RRRPPYLP RPP 15
; Db 279 RRPPIQLRPP 292
;
; RESULT 15
; US-09-253-682-26
; Sequence 26, Application US/09253682
; Patent No. 6040170
; GENERAL INFORMATION:
; APPLICANT: Spaete, Richard
; APPLICANT: Cha, Tai-An
; TITLE OF INVENTION: NOVEL HUMAN CYTOMEGALOVIRUS
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Luann Cserr Attorney at Law
; STREET: 750 Arimo Avenue
; CITY: Oakland
; STATE: CA
; COUNTRY: USA
; ZIP: 94610
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/253,682
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/926,922
; FILING DATE: September 10, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Cserr, Luann
; REGISTRATION NUMBER: 31,822
; REFERENCE/DOCKET NUMBER: AVIR 11A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 510-834-1448
; TELEFAX: 510-839-7810
; INFORMATION FOR SEQ ID NO: 26:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 336 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein

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Db 279 RREIPFLOPRPP 292

## RESULT 15

PCT-US95-12080-3  
; Sequence 3, Application PC/TUS9512080  
; GENERAL INFORMATION:  
; APPLICANT: Children's Medical Center Corporation  
; TITLE OF INVENTION: Synducin Mediated Modulation of Tissue Repair  
; NUMBER OF SEQUENCES: 4  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Patrea L. Pabet  
; STREET: 2800 One Atlantic Center  
; STREET: 1201 West Peachtree  
; CITY: Atlanta  
; STATE: Georgia  
; COUNTRY: USA  
; ZIP: 30309-3450  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent in Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: PCT/US95/12080  
; FILING DATE:  
; CLASSIFICATION:  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (404)-873-8794  
; TELEFAX: (404)-815-8795  
; INFORMATION FOR SEQ ID NO: 3:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 59 amino acids  
; TYPE: amino acid  
; TOPOLOGY: linear  
; PCT-US95-12080-3

Query Match 65.6%; Score 59; DB 5; Length 59;  
Best Local Similarity 84.6%; Pred. No. 0.57;  
Matches 11; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPYLP RPR 13  
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Db 2 RIRPRPRLPRR 14

Search completed: May 13, 2003, 10:42:08  
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GenCore version 5.1.4 p5 4578  
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:40:37 ; Search time 17 Seconds  
(without alignments)  
81.199 Million cell updates/sec

Title: US-09-426-011D-3

Perfect score: 90

Sequence: 1 RRRPPPLPRPRP 15

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Searched: 349150 seqs, 92025710 residues

Total number of hits satisfying chosen parameters: 349150

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Published Applications AA:\*

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- 2: /cgn2\_6/ptodata/2/pubpaa/PCT NEW PUB.pap:\*
- 3: /cgn2\_6/ptodata/2/pubpaa/US05 NEW PUB.pap:\*
- 4: /cgn2\_6/ptodata/2/pubpaa/US06\_PUBCOMB.pap:\*
- 5: /cgn2\_6/ptodata/2/pubpaa/US07 NEW PUB.pap:\*
- 6: /cgn2\_6/ptodata/2/pubpaa/US07\_PUBCOMB.pap:\*
- 7: /cgn2\_6/ptodata/2/pubpaa/PCTUS\_PUBCOMB.pap:\*
- 8: /cgn2\_6/ptodata/2/pubpaa/US08\_PUBCOMB.pap:\*
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- 10: /cgn2\_6/ptodata/2/pubpaa/US09\_PUBCOMB.pap:\*
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- 12: /cgn2\_6/ptodata/2/pubpaa/US10\_PUBCOMB.pap:\*
- 13: /cgn2\_6/ptodata/2/pubpaa/US60 NEW PUB.pap:\*
- 14: /cgn2\_6/ptodata/2/pubpaa/US60\_PUBCOMB.pap:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	66	73.3	59	10	US-09-030-619-163
2	60	66.7	953	10	US-09-888-615-66
3	56.5	62.8	74	10	US-09-864-761-45555
4	55	61.1	45	10	US-09-864-761-49065
5	54	60.0	250	9	US-10-102-806-517
6	53	58.9	18	10	US-09-030-619-96
7	53	58.9	18	10	US-09-030-619-158
8	53	58.9	18	10	US-09-030-619-159
9	52	57.8	18	10	US-09-030-619-160
10	52	57.8	354	9	US-10-004-717-2
11	52	57.8	354	9	US-10-004-717-58
12	51	56.7	180	10	US-09-997-701-5
13	51	56.7	195	10	US-09-997-701-1
14	50.5	56.1	392	10	US-09-747-835A-55
15	50.5	56.1	393	9	US-10-243-035-2
16	50.5	56.1	419	10	US-09-828-035-2
17	50.5	56.1	1314	10	US-09-747-835A-29
18	50	55.6	99	10	US-09-864-761-43778
19	50	55.6	146	9	US-09-989-920-237

20	50	55.6	449	9	US-10-125-540-320	Sequence 320, App
21	50	55.6	449	9	US-10-103-313-438	Sequence 438, App
22	50	55.6	449	10	US-09-764-870-320	Sequence 320, App
23	50	55.6	449	10	US-09-764-853-643	Sequence 643, App
24	49.5	55.0	129	9	US-09-975-719-97	Sequence 97, Appl
25	49	54.4	223	12	US-10-062-254-204	Sequence 204, App
26	48.5	53.9	111	10	US-09-864-761-47005	Sequence 47005, A
27	48	53.3	281	8	US-08-971-317A-6	Sequence 6, Appli
28	48	53.3	281	9	US-09-131-237-6	Sequence 6, Appli
29	48	53.3	281	9	US-10-174-554-10	Sequence 10, Appl
30	48	53.3	281	9	US-10-151-882-44	Sequence 44, Appl
31	48	53.3	281	10	US-09-802-669-25	Sequence 25, Appl
32	48	53.3	281	10	US-09-193-663-6	Sequence 6, Appli
33	48	53.3	281	10	US-09-027-287-6	Sequence 6, Appli
34	48	53.3	281	10	US-09-252-656B-6	Sequence 6, Appli
35	48	53.3	281	10	US-09-929-493-6	Sequence 6, Appli
36	48	53.3	281	10	US-09-927-110-1	Sequence 1, Appli
37	48	53.3	281	12	US-10-012-452-13	Sequence 13, Appl
38	48	53.3	1134	9	US-10-001-873-50	Sequence 50, Appl
39	47.5	52.8	272	10	US-09-925-300-1697	Sequence 1697, Ap
40	47	52.2	42	10	US-09-030-619-162	Sequence 162, App
41	47	52.2	84	9	US-10-174-590-486	Sequence 486, App
42	47	52.2	84	9	US-10-176-758-486	Sequence 486, App
43	47	52.2	84	9	US-10-175-737-486	Sequence 486, App
44	47	52.2	84	9	US-10-173-706-486	Sequence 486, App
45	47	52.2	84	9	US-10-175-738-486	Sequence 486, App

ALIGNMENTS

RESULT 1

US-09-030-619-163

; Sequence 163, Application US/09030619B

; Patent No. US20020035061A1

; GENERAL INFORMATION:

; APPLICANT: Krieger, Timothy J.

; APPLICANT: Taylor, Robert

; APPLICANT: Erfile, Douglas

; APPLICANT: Fraser, Janet R.

; APPLICANT: West, Michael H.P.

; APPLICANT: McNicol, Patricia J.

; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING

; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION

; TITLE OF INVENTION: WITH ANTIBIOTICS

; FILE REFERENCE: 66081.406

; CURRENT APPLICATION NUMBER: US/09/030,619B

; CURRENT FILING DATE: 1998-02-25

; NUMBER OF SEQ ID NOS: 232

; SOFTWARE: FastSeq for Windows Version 3.0

; SEQ ID NO 163

; LENGTH: 59

; TYPE: PRT

; ORGANISM: Bos taurus

US-09-030-619-163

Query Match 73.3%; Score 66; DB 10; Length 59;

Best Local Similarity 85.7%; Pred. No. 0.27; Mismatches 2; Indels 0; Gaps 0;

Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 RRRPPPLPRPRP 14

| | | | | | | | | | | | | |

Db 2 RRRPPPLPRPRP 15

| | | | | | | | | | | | | |

RESULT 2

US-09-888-615-66

; Sequence 66, Application US/09888615

; Patent No. US20020064856A1

; GENERAL INFORMATION:

; APPLICANT: PLOWMAN, GREGORY

; APPLICANT: WHYTE, DAVID

; APPLICANT: CAENEPEEL, SEAN



; PRIOR APPLICATION NUMBER: US 60/234,687  
; PRIOR FILING DATE: 2000-09-21  
; PRIOR APPLICATION NUMBER: US 09/608,408  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: US 09/774,203  
; PRIOR FILING DATE: 2001-01-29  
; NUMBER OF SEQ ID NOS: 49117  
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 49065  
; LENGTH: 45  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AC005973.2  
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 0.88  
; OTHER INFORMATION: EST\_HUMAN HIT: A1358103.1, EVALUATE 4.60e+00  
US-09-864-761-49065

Query Match 61.1%; Score 55; DB 10; Length 45;  
Best Local Similarity 76.9%; Pred. No. 3.6;  
Matches 10; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
QY 1 RRRPRPPLPRPR 13  
DB 19 RRRPRPPGPRPP 31

RESULT 5  
US-10-102-806-517  
; Sequence 517, Application US/10102806  
; Publication No. US20030054421A1  
; GENERAL INFORMATION:  
; APPLICANT: Rosen et al.  
; TITLE OF INVENTION: Nucleic Acids, Proteins and Antibodies  
; FILE REFERENCE: PAM03P1C1  
; CURRENT APPLICATION NUMBER: US/10/102,806  
; PRIOR FILING DATE: 2002-03-22  
; PRIOR APPLICATION NUMBER: 09/925,298  
; PRIOR FILING DATE: 2001-08-10  
; PRIOR APPLICATION NUMBER: PCT/US00/05881  
; PRIOR FILING DATE: 2000-03-08  
; PRIOR APPLICATION NUMBER: 60/124,270  
; PRIOR FILING DATE: 1999-03-12  
; NUMBER OF SEQ ID NOS: 846  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 517  
; LENGTH: 250  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: SITE  
; LOCATION: (118)  
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids  
; NAME/KEY: SITE  
; LOCATION: (161)  
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids  
; NAME/KEY: SITE  
; LOCATION: (204)  
; OTHER INFORMATION: Xaa equals any of the naturally occurring L-amino acids  
US-10-102-806-517

Query Match 60.0%; Score 54; DB 9; Length 250;  
Best Local Similarity 71.4%; Pred. No. 23;  
Matches 10; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 RRRPRPPLPRPR 14  
DB 202 RXHRPPAAPRPP 215

RESULT 6  
US-09-030-619-96  
; Sequence 96, Application US/09030619B

; Patent No. US20020035061A1  
; GENERAL INFORMATION:  
; APPLICANT: Krieger, Timothy J.  
; APPLICANT: Taylor, Robert  
; APPLICANT: Erfile, Douglas  
; APPLICANT: Fraser, Janet R.  
; APPLICANT: West, Michael H.P.  
; APPLICANT: McNicol, Patricia J.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING  
; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION  
; TITLE OF INVENTION: WITH ANTIBIOTICS  
; FILE REFERENCE: 660081.406  
; CURRENT APPLICATION NUMBER: US/09/030,619B  
; CURRENT FILING DATE: 1998-02-25  
; NUMBER OF SEQ ID NOS: 232  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 96  
; LENGTH: 18  
; TYPE: PRT  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Cationic Peptide Analogue  
US-09-030-619-96

Query Match 58.9%; Score 53; DB 10; Length 18;  
Best Local Similarity 72.7%; Pred. No. 2.6;  
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
QY 5 RPPYLPFRPP 15  
DB 4 RPYVIFQPRPP 14

RESULT 7  
US-09-030-619-158  
; Sequence 158, Application US/09030619B  
; Patent No. US20020035061A1  
; GENERAL INFORMATION:  
; APPLICANT: Krieger, Timothy J.  
; APPLICANT: Taylor, Robert  
; APPLICANT: Erfile, Douglas  
; APPLICANT: Fraser, Janet R.  
; APPLICANT: West, Michael H.P.  
; APPLICANT: McNicol, Patricia J.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING  
; TITLE OF INVENTION: INFECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION  
; TITLE OF INVENTION: WITH ANTIBIOTICS  
; FILE REFERENCE: 660081.406  
; CURRENT APPLICATION NUMBER: US/09/030,619B  
; CURRENT FILING DATE: 1998-02-25  
; NUMBER OF SEQ ID NOS: 232  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 158  
; LENGTH: 18  
; TYPE: PRT  
; ORGANISM: Apis mellifera  
US-09-030-619-158

Query Match 58.9%; Score 53; DB 10; Length 18;  
Best Local Similarity 72.7%; Pred. No. 2.6;  
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
QY 5 RPPYLPFRPP 15  
DB 4 RPYVIFQPRPP 14

RESULT 8  
US-09-030-619-159  
; Sequence 159, Application US/09030619B  
; Patent No. US20020035061A1  
; GENERAL INFORMATION:  
; APPLICANT: Krieger, Timothy J.

```
; APPLICANT: Taylor, Robert
; APPLICANT: Exfle, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INJECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTIBIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030.619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 159
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Apis mellifera
US-09-030-619-159

Query Match      58.9%; Score 53; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 2.6;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYLPRPRPP 15
   |||:||||
Db 4 RPYIQPRPP 14

RESULT 9
US-09-030-619-160
; Sequence 160, Application US/09030619B
; Patent No. US20020035061A1
; GENERAL INFORMATION:
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Exfle, Douglas
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: McNicol, Patricia J.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INJECTIONS USING CATIONIC PEPTIDES ALONE OR IN COMBINATION
; TITLE OF INVENTION: WITH ANTIBIOTICS
; FILE REFERENCE: 660081.406
; CURRENT APPLICATION NUMBER: US/09/030.619B
; CURRENT FILING DATE: 1998-02-25
; NUMBER OF SEQ ID NOS: 232
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 160
; LENGTH: 18
; TYPE: PRT
; ORGANISM: Apis mellifera
US-09-030-619-160

Query Match      57.8%; Score 52; DB 10; Length 18;
Best Local Similarity 72.7%; Pred. No. 3.3;
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYLPRPRPP 15
   |||:||||
Db 4 RPYIQPRPP 14

RESULT 10
US-10-004-717-2
; Sequence 2, Application US/10004717
; Publication No. US20020192665A1
; GENERAL INFORMATION:
; APPLICANT: ZOGHBI, HUDA Y.
; APPLICANT: YANG, QI
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPEUTIC USE OF AN
; TITLE OF INVENTION: ATONAL ASSOCIATED SEQUENCE FOR DEAFNESS,
; TITLE OF INVENTION: OSTEOARTHRITIS, AND ABNORMAL CELL PROLIFERATION
; FILE REFERENCE: P01899US4
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; CURRENT APPLICATION NUMBER: US/10/004,717
; CURRENT FILING DATE: 2002-08-16
; PRIOR APPLICATION NUMBER: 09/585,645
; PRIOR FILING DATE: 2000-06-01
; PRIOR APPLICATION NUMBER: 60/176,993
; PRIOR FILING DATE: 2000-01-19
; PRIOR APPLICATION NUMBER: 60/137,060
; PRIOR FILING DATE: 1999-06-01
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 2
; LENGTH: 354
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-004-717-2

Query Match      57.8%; Score 52; DB 9; Length 354;
Best Local Similarity 57.1%; Pred. No. 52;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRPYPYLPYRPP 15
   |||:||||
Db 21 RQPHHLPQPP 34

RESULT 11
US-10-004-717-58
; Sequence 58, Application US/10004717
; Publication No. US20020192665A1
; GENERAL INFORMATION:
; APPLICANT: ZOGHBI, HUDA Y.
; APPLICANT: YANG, QI
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPEUTIC USE OF AN
; TITLE OF INVENTION: ATONAL ASSOCIATED SEQUENCE FOR DEAFNESS,
; TITLE OF INVENTION: OSTEOARTHRITIS, AND ABNORMAL CELL PROLIFERATION
; FILE REFERENCE: P01899US4
; CURRENT APPLICATION NUMBER: US/10/004,717
; CURRENT FILING DATE: 2002-08-16
; PRIOR APPLICATION NUMBER: 09/585,645
; PRIOR FILING DATE: 2000-06-01
; PRIOR APPLICATION NUMBER: 60/176,993
; PRIOR FILING DATE: 2000-01-19
; PRIOR APPLICATION NUMBER: 60/137,060
; PRIOR FILING DATE: 1999-06-01
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 58
; LENGTH: 354
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-004-717-58

Query Match      57.8%; Score 52; DB 9; Length 354;
Best Local Similarity 57.1%; Pred. No. 52;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRPYPYLPYRPP 15
   |||:||||
Db 21 RQPHHLPQPP 34

RESULT 12
US-09-997-701-5
; Sequence 5, Application US/09997701
; Patent No. US20020107180A1
; GENERAL INFORMATION:
; APPLICANT: Yue, Henry
; APPLICANT: Corley, Neil C.
; APPLICANT: Guesler, Karl J.
; APPLICANT: Gorgone, Gina A.
; APPLICANT: Baughn, Mariah R.
; TITLE OF INVENTION: CELL SURFACE GLYCOPROTEINS
; FILE REFERENCE: PF-0631 US
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; CURRENT APPLICATION NUMBER: US/09/997,701  
; CURRENT FILING DATE: 2001-11-30  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/470,946  
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-12-22  
; NUMBER OF SEQ ID NOS: 6  
; SOFTWARE: PERL Program  
; SEQ ID NO 5  
; LENGTH: 180  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: g2499136  
US-09-997-701-5

Query Match 56.7%; Score 51; DB 10; Length 180;  
Best Local Similarity 53.8%; Pred. No. 36;  
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRRPPYLP-RRPP 15  
:|:|:|:|:|:|  
Db 47 KRPYPYQENP 59

## RESULT 13

US-09-997-701-1  
; Sequence 1, Application US/09997701  
; Patent No. US20020107180A1  
; GENERAL INFORMATION:  
; APPLICANT: Yue, Henry  
; APPLICANT: Corley, Neil C.  
; APPLICANT: Guegler, Karl J.  
; APPLICANT: Gorgone, Gina A.  
; APPLICANT: Baughn, Mariah R.  
; TITLE OF INVENTION: CELL SURFACE GLYCOPROTEINS  
; FILE REFERENCE: PF-0631 US  
; CURRENT APPLICATION NUMBER: US/09/997,701  
; CURRENT FILING DATE: 2001-11-30  
; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/470,946  
; PRIOR FILING DATE: EARLIER FILING DATE: 1999-12-22  
; NUMBER OF SEQ ID NOS: 6  
; SOFTWARE: PERL Program  
; SEQ ID NO 1  
; LENGTH: 195  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: 2297891  
US-09-997-701-1

Query Match 56.7%; Score 51; DB 10; Length 195;  
Best Local Similarity 53.8%; Pred. No. 39;  
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRRPPYLP-RRPP 15  
:|:|:|:|:|:|  
Db 47 KRPYPYQENP 59

## RESULT 14

US-09-747-835A-55  
; Sequence 55, Application US/09747835A  
; Patent No. US20020146692A1  
; GENERAL INFORMATION:  
; APPLICANT: Yamazaki, Victoria  
; APPLICANT: Tang, Y. Tom  
; APPLICANT: Liu, Chinghua  
; APPLICANT: Zhou, Ping  
; APPLICANT: Wang, Dunrui  
; APPLICANT: Zhang, Jie  
; APPLICANT: Ren, Feiyan  
; APPLICANT: Asundi, Vinod  
; APPLICANT: Drmanac, Radoje T  
; TITLE OF INVENTION: METHODS AND MATERIALS RELATING TO G PROTEIN-COUPLED RECEPTOR-LIKE

; TITLE OF INVENTION: LIKE) POLYPEPTIDES AND POLYNUCLEOTIDES  
; FILE REFERENCE: HYS-37CIP  
; CURRENT APPLICATION NUMBER: US/09/747,835A  
; CURRENT FILING DATE: 2002-03-08  
; PRIOR APPLICATION NUMBER: US 09/729,739  
; PRIOR FILING DATE: 2000-12-04  
; PRIOR APPLICATION NUMBER: US 09/653,450  
; PRIOR FILING DATE: 2000-08-31  
; PRIOR APPLICATION NUMBER: US 09/620,312  
; PRIOR FILING DATE: 2000-07-19  
; PRIOR APPLICATION NUMBER: US 09/598,042  
; PRIOR FILING DATE: 2000-06-20  
; PRIOR APPLICATION NUMBER: US 09/552,317  
; PRIOR FILING DATE: 2000-04-25  
; PRIOR APPLICATION NUMBER: US 09/488,725  
; PRIOR FILING DATE: 2000-01-21  
; NUMBER OF SEQ ID NOS: 63  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 55  
; LENGTH: 392  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
US-09-747-835A-55

Query Match 56.1%; Score 50.5; DB 10; Length 392;  
Best Local Similarity 68.8%; Pred. No. 85;  
Matches 11; Conservative 0; Mismatches 4; Indels 1; Gaps 1;

QY 1 RRRPPYLP-RRPP 15  
|:|:|:|:|:|:|  
Db 368 RRRPNPVRPRGP 383

## RESULT 15

US-10-243-035-2  
; Sequence 2, Application US/10243035  
; Publication No. US20030049697A1  
; GENERAL INFORMATION:  
; APPLICANT: LAZDUNSKI, MICHEL  
; APPLICANT: LESAGE, FLORIAN  
; TITLE OF INVENTION: NEW FAMILY OF MECHANOSENSITIVE HUMAN POTASSIUM CHANNELS  
; TITLE OF INVENTION: ACTIVATED BY POLYUNSATURATED FATTY ACIDS AND THEIR USE  
; FILE REFERENCE: 1317-02  
; CURRENT APPLICATION NUMBER: US/10/243,035  
; CURRENT FILING DATE: 2002-09-13  
; NUMBER OF SEQ ID NOS: 15  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 2  
; LENGTH: 393  
; TYPE: PRT  
; ORGANISM: Homo sapiens  
US-10-243-035-2

Query Match 56.1%; Score 50.5; DB 9; Length 393;  
Best Local Similarity 68.8%; Pred. No. 85;  
Matches 11; Conservative 0; Mismatches 4; Indels 1; Gaps 1;

QY 1 RRRPPYLP-RRPP 15  
|:|:|:|:|:|:|  
Db 368 RRRPNPVRPRGP 383

Search completed: May 13, 2003, 10:42:32  
Job time : 18 secs

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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:38:17 ; Search time 16 seconds  
(without alignments)  
90.126 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPPPYLPRPP 15

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 73:\*

1: pir1:\*

2: pir2:\*

3: pir3:\*

4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	90	100.0	172	2	S68232 antimicrobial prot
2	66	73.3	59	2	A36589 bacterinec 7 - bov
3	59.5	66.1	82	2	A41051 spore coat protein
4	58	64.4	190	2	S68230 antimicrobial pept
5	56.5	62.8	168	2	S35330 apidaecin 14 precu
6	56.5	62.8	199	2	S14981 extensin class I (
7	55	61.1	437	2	A88942 protein R13D11.3
8	54.5	60.6	301	2	JQ1563 hybrid proline-ric
9	54	60.0	359	2	T13478 hypothetical prote
10	54	60.0	437	2	T32652 hypothetical prote
11	53	58.9	26	2	S06675 apidaecin 1b precu
12	53	58.9	144	2	S35331 apidaecin 22 precu
13	53	58.9	184	2	T29373 hypothetical prote
14	53	58.9	283	2	S35332 apidaecin 73 precu
15	53	58.9	428	2	E71415 probable coll wall
16	53	58.9	491	2	T07598 proline-rich prot
17	52	57.8	261	1	WMBEXE infected cell prot
18	52	57.8	439	2	S51939 chitinase (EC 3.2
19	52	57.8	467	2	S71169 protein kinase, 54
20	52	57.8	1066	2	G86292 hypothetical prote
21	51.5	57.2	1187	1	JC4155 protein-tyrosine-p
22	51.5	57.2	1189	1	JC2366 protein-tyrosine-p
23	51	56.7	180	2	S43791 PBDX protein - hum
24	50.5	56.1	1216	2	JW0105 synaptotagmin 2 alp
25	50	55.6	192	2	S76867 hypothetical prote
26	50	55.6	383	2	T06753 zinc finger protei
27	50	55.6	415	1	A34170 acrosin (EC 3.4.21
28	50	55.6	421	2	S29599 acrosin (EC 3.4.21
29	50	55.6	424	2	A54964 spliceosome-associ

30	50	55.6	449	2	S16748	proline-rich prote
31	50	55.6	547	2	C96828	unknown protein F1
32	50	55.6	1460	1	EDBEIF	immediate-early pr
33	50	55.6	3036	2	T18995	hypothetical prote
34	49.5	55.0	589	2	T29299	hypothetical prote
35	49	54.4	118	2	T19345	hypothetical prote
36	49	54.4	134	2	JC5572	proline-rich prote
37	49	54.4	161	2	F72593	hypothetical prote
38	49	54.4	210	2	T33700	hypothetical prote
39	49	54.4	218	2	T22261	hypothetical prote
40	49	54.4	296	2	A27319	gliadin - wheat
41	49	54.4	296	2	S07361	alpha/beta-gliadin
42	49	54.4	352	2	F84799	hypothetical prote
43	49	54.4	369	2	S20500	hydroxyproline-ric
44	49	54.4	380	2	T32944	hypothetical prote
45	49	54.4	413	2	H87604	hypothetical prote

ALIGNMENTS

RESULT 1

S68232

antimicrobial protein PR-39 precursor, cathelin-associated - pig

N:Alternate names: myeloid antibacterial protein PR-39

C:Species: Sus scrofa domestica (domestic pig)

C>Date: 15-Feb-1997 #sequence revision 13-Mar-1997 #text\_change 20-Jun-2000

C:Accession: S68232, JN0899; I47138; S19563

R:Zhao, C.; Ganz, T.; Lehrer, R.I.

FEBS Lett. 376, 130-134, 1995

A:Title: Structures of genes for two cathelin-associated antimicrobial peptides: propheni

A:Reference number: S68232; MUID:96105365; PMID:7498526

A:Accession: S68232

A>Status: translation not shown

A:Molecule type: DNA

A:Residues: 1-172 <ZHA>

A:Cross-references: EMBL:X89201; NID:g1165150; PIDN:CAA61487.1; PID:g1165151

A:Experimental source: leukocytes

R:Storici, P.; Zanetti, M.

Biochem. Biophys. Res. Commun. 196, 1058-1065, 1993

A:Title: A cDNA derived from pig bone marrow cells predicts a sequence identical to the

A:Reference number: JN0899; MUID:94071853; PMID:8250863

A:Accession: JN0899

A:Molecule type: mRNA

A:Residues: 1-20, 'A', 22-172 <STO>

A:Cross-references: GB:L23825; NID:9435100; PIDN:AAA31109.1; PID:9435101

A:Experimental source: bone marrow cells

R:Gudmundsson, G.H.; Magnusson, K.P.; Chowdhary, B.P.; Johansson, M.; Andersson, L.; Bom

Proc. Natl. Acad. Sci. U.S.A. 92, 7085-7089, 1995

A:Title: Structure of the gene for porcine peptide antibiotic PR-39, a cathelin gene fami

A:Reference number: I47138; MUID:95350216; PMID:7624374

A:Accession: I47138

A>Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-28, 'T', 30-89, 'QR', 92-116, 'NDP', 120-172 <GUD>

A:Cross-references: EMBL:X87236; NID:9829142; PIDN:CAA60682.1; PID:g1051298

R:Ageberth, B.; Lee, J.Y.; Bergman, T.; Carlquist, M.; Boman, H.G.; Mutt, V.; Joernvall,

Eur. J. Biochem. 202, 849-854, 1991

A:Title: Amino acid sequence of PR-39. Isolation from pig intestine of a new member of t

A:Reference number: S19563; MUID:92111534; PMID:1765098

A:Accession: S19563

A:Molecule type: protein

A:Residues: 131-169 <AGE>

A:Experimental source: intestine

C:Genetics:

A:Gene: PR39

C:Superfamily: cathelin; cystatin homology

C:Keywords: amidated carboxyl end; antibacterial

F:1-29/Domain: signal sequence #status predicted <SIG>

F:22-129/Domain: cystatin homology <CYS>

F:30-130/Domain: propeptide #status predicted <PRO>

F:131-169/Product: antimicrobial protein PR-39 #status experimental <MAT>

F;169/Modified site: amidated carboxyl end (Pro) (amide in mature form from following gl

Query Match 100.0%; Score 90; DB 2; Length 172;  
 Best Local Similarity 100.0%; Pred. No. 0.00035;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 15  
 |||||  
 DB 131 RRRPRPPVLPRLPRPP 145

RESULT 2  
 A36589  
 bacitracin 7 - bovine  
 C;Species: Bos primigenius taurus (cattle)  
 C;Date: 12-Apr-1991 #sequence\_revision 12-Apr-1991 #text\_change 09-May-1997  
 C;Accession: A36589  
 R;Frank, R.W.; Gennaro, R.; Schneider, K.; Przybylski, M.; Romeo, D.  
 J. Biol. Chem. 265, 18871-18874, 1990  
 A;Title: Amino acid sequences of two proline-rich bacitracins. Antimicrobial peptides of  
 A;Reference number: A36589; MUID:91035404; PMID:2229048  
 A;Accession: A36589  
 A;Status: preliminary  
 A;Molecule type: protein  
 A;Residues: 1-59 <PRA>  
 C;Superfamily: cathelin; cystatin homology

Query Match 73.3%; Score 66; DB 2; Length 59;  
 Best Local Similarity 85.7%; Pred. No. 0.086;  
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 14  
 |||||  
 DB 2:RIRPRPRLPRPP 15

RESULT 3  
 A41051  
 spore coat protein precursor - Bacillus subtilis  
 C;Species: Bacillus subtilis  
 C;Date: 03-Apr-1992 #sequence\_revision 06-Jan-1995 #text\_change 11-Jan-2002  
 C;Accession: S04835; A41051; F69606  
 R;Aranson, A.I.; Song, H.Y.; Bourne, N.  
 Mol. Microbiol. 3, 437-444, 1989  
 A;Title: Gene structure and precursor processing of a novel Bacillus subtilis spore coat  
 A;Reference number: S04835; MUID:89313296; PMID:2546006  
 A;Accession: S04835  
 A;Molecule type: DNA  
 A;Residues: 'MNVHTPNLSIRNMVGIKAREVFL', 2-82 <AR2>  
 A;Cross-references: EMBL:X13740; NID:G39864; PIDN:CAA32004.1; PID:G39865  
 A;Experimental source: strain JH642  
 A;Note: Part of this sequence, including the amino end of the mature protein, was confir  
 R;Bourne, N.; FitzJames, P.C.; Aranson, A.I.  
 J. Bacteriol. 173, 6618-6625, 1991  
 A;Title: Structural and germination defects of Bacillus subtilis spores with altered con  
 A;Reference number: A41051; MUID:92011439; PMID:1917883  
 A;Accession: A41051  
 A;Molecule type: protein  
 A;Residues: 'XX', 3-11 <BOU>  
 A;Experimental source: strain JH642  
 A;Note: The material sequenced was the larger of two isolated precursor forms; the amin  
 A;Note: both the location of the transcription start site and peptide sequencing of the  
 R;Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Berter  
 C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Chd  
 A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.  
 Nature 390, 249-256, 1997  
 A;Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Galizzi, A.; Gallen  
 iech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holsappel, S.; Hosono, S.; Hullo, M.F.  
 Koetter, P.; Konigstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapius, A.; Lardinois  
 A;Authors: Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maueel  
 Y, M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portetelle  
 Rieger, M.; Rivolta, C.; Rocha, E.; Roche, M.; Rose, M.; Sadaie, Y.; Sato, T.; Scanlon,  
 A;Authors: Schleich, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seron

akeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Tosato, V.; Uchiyama,  
 T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, K  
 A;Authors: Yoshikawa, H.F.; Zumstein, E.; Yoshikawa, H.; Danchin, A.  
 A;Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.  
 A;Reference number: A69580; MUID:98044033; PMID:9384377  
 A;Accession: F69606  
 A;Status: preliminary; nucleic acid sequence not shown; translation not shown  
 A;Molecule type: DNA  
 A;Residues: 'MNVHTPNLSIRNMVGIKAREVFL', 2-82 <KUN>  
 A;Cross-references: GB:Z99110; GB:AL009126; NID:G2633472; PIDN:CAB13066.1; PID:G2633563  
 A;Experimental source: strain 168  
 C;Comment: This structural protein is expressed during stage V of sporulation.  
 C;Genetics:  
 A;Gene: cotT  
 A;Start codon: TTG  
 C;Keywords: sporulation  
 F;1-19/Domin: propeptide #status experimental <PRO>  
 F;20-82/Product: spore coat protein #status experimental <MAT>

Query Match 66.1%; Score 59.5; DB 2; Length 82;  
 Best Local Similarity 84.6%; Pred. No. 0.69;  
 Matches 11; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 4 PRPP-YLPRLPRPP 15  
 |||||  
 DB 49 PRPPYYPRPP 61

RESULT 4  
 S68230  
 antimicrobial peptide precursor - sheep  
 N;Alternate names: Bac7.5 peptide homolog  
 C;Species: Ovis orientalis aries, Ovis ammon aries (domestic sheep)  
 C;Date: 15-Feb-1997 #sequence\_revision 13-Mar-1997 #text\_change 21-Jan-2000  
 C;Accession: S68230  
 R;Bagella, L.; Scocchi, M.; Zanetti, M.  
 FEBS Lett. 376, 225-228, 1995  
 A;Title: cDNA sequences of three sheep myeloid cathelicidins.  
 A;Reference number: S68228; MUID:96105386; PMID:7498547  
 A;Accession: S68230  
 A;Status: preliminary  
 A;Molecule type: mRNA  
 A;Residues: 1-190 <BAG>  
 A;Cross-references: EMBL:L46852; NID:g1161244; PIDN:AAA85468.1; PID:g1161245  
 C;Superfamily: cathelin; cystatin homology  
 F;1-23/Domin: signal sequence #status predicted <SIG>  
 F;22-129/Domin: signal sequence #status predicted <SIG>  
 F;29-130/Domin: cystatin homology <CYS>  
 F;130-190/Product: propeptide #status predicted <PRO>  
 F;130-190/Product: antimicrobial peptide #status predicted <MAT>

Query Match 64.4%; Score 58; DB 2; Length 190;  
 Best Local Similarity 78.6%; Pred. No. 2.4;  
 Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RRRPRPPVLPRLPRPP 14  
 |||||  
 DB 132 RLRPRPRLPRPP 145

RESULT 5  
 S35330  
 apidaecin 14 precursor - honeybee  
 N;Contains: apidaecin II  
 C;Species: Apis mellifera (honeybee)  
 C;Date: 03-Feb-1994 #sequence\_revision 03-Feb-1994 #text\_change 21-Jul-2000  
 C;Accession: S35330; S06676  
 R;Casteels-Josson, K.; Capaci, T.; Casteels, P.; Tempst, P.  
 EMBO J. 12, 1569-1578, 1993  
 A;Title: Apidaecin multipetide precursor structure: a putative mechanism for amplificati  
 A;Reference number: S35330; MUID:93223697; PMID:8467807  
 A;Accession: S35330  
 A;Molecule type: mRNA  
 A;Residues: 1-168 <CAS>

A; Cross-references: EMBL:X72575; NID:g297062; PIDN:CAA51167.1; PID:g297063  
R; Casteels, P.; Ampe, C.; Jacobs, F.; Vaeck, M.; Tempst, P.  
EMBO J. 8, 2387-2391, 1989  
A; title: Apidaecins: antibacterial peptides from honeybees.  
A; Reference number: S05383; MUID:90005446; PMID:2676519

A;Accession: S06676  
A;Molecule type: protein  
A;Residues: 43-60 <CA2>  
C;Superfamily: procylic acid repetitive protein  
F;43-60/Product: apidaecin II #status experimental <MAT>

Query Match	62.8%	Score	56.5	DB 2	Length	168	.
Best Local Similarity	50.0%	Pred. No.	3.2				
Matches	11	Conservative	2	Mismatches	2	Indels	7
						Gaps	1
							1

Qy 1 RRRP-----RPPYLP RPPPP 15  
||| ||| ||| :|||  
Db 117 RREPEAEFGNNRPVYIOP RPP 138

RESULT 6  
S14981  
extensin class I (clone w1-8 L) - tomato (fragment)  
C:Species: Lycopersicon esculentum (tomato)  
C:date: 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change 11-Jan-2000  
C:Accession: S14981  
R:Showalter, A.M.; Zhou, J.; Rumsau, D.; Worst, S.G.; Varner, J.E.

Plant Mol. Biol. 16, 547-565, 1991  
A;Title: Tomato extensin and extensin-like cDNAs: structure and expression in response  
A;Reference number: S14970; MUID:91329690; PMID:1714316

A;Accession: S14981  
A;Status: preliminary  
A;Molecule type: mRNA  
A;Residues: 1-199 <SHO>  
A;Cross-references: EMBL:X55692  
A;Experimental source: cv. UC82B  
C;Superfamily: hydroxyproline-rich glycoprotein  
C;Keywords: Cell wall, glycoprotein, hydroxyproline

Query Match	62.8%	Score	56.5	DB 2	Length	199			
Best Local Similarity	73.3%	Pred. No.	3.7						
Matches	11	Conservative	0	Mismatches	1	Indels	3	Gaps	1

Qy 4 PRPP---YLPRRPP 15  
Db 77 PRPPPEYLPPEPRPP 91

RESULT 7  
A88942 protein R13D11.3 [imported] - *Caenorhabditis elegans*  
C;Species: *Caenorhabditis elegans*  
C;Date: 10-May-2001 #sequence\_revision 10-May-2001 #text\_change 10-May-2001  
C;Accession: A88942  
R;anonymous, The C. elegans Sequencing Consortium.  
Science 282, 2012-2018, 1998  
A;Title: Genome sequence of the nematode *C. elegans*: a platform for investigating biological processes  
A;Reference number: A75000; PMID:9851916  
A;Note: see websites genome.wustl.edu/gsc/C.elegans/ and www.sanger.ac.uk/Projects/C.elegans/ for more information  
A;Note: published errata appeared in Science 283, 35, 1999; Science 283, 2103, 1999; and Science 283, 2104, 1999.  
A;Accession: A88942  
A;Status: preliminary  
A;Molecule type: DNA  
A;Residues: 1-437 <STO>  
A;Cross-references: GB:chr\_V; PIDN:AAB69949.1; PID:g2384928; GSPDB:GN00023; CESP:R13D11.3  
C;Genetics:  
A;Gene: R13D11.3  
A;Map position: 5

Query Match	61.1%	Score 55;	DB 2;	Length 437;
Best Local Similarity	75.0%	Pred. No. 12;		
Matches	9;	Conservative	1;	Mismatches
			2;	Indels
			0;	Gaps

Qy 4 PRPPYLP RPPRP 15  
|||: ||||  
Db 23 PRPPHPP IPRPP 34

## RESULT 8

JQ1663  
 hybrid proline-rich protein - maize  
 C:Species: Zea mays (maize)  
 C:Date: 30-Sep-1993 #sequence\_revision 30-Sep-1993 #text\_change 24-Sep-1999  
 C:Accession: JQ1663  
 P:Jose-Estanyol, M.; Ruiz-Avila, L.; Puigdomenech, P.  
 Plant Cell 4, 413-423, 1992  
 A:Title: A maize embryo-specific gene encodes a proline-rich and hydrophobic protein.  
 A:Reference number: JQ1663; MUID:92361259; PMID:1498500

## A;Accession: JQ1663

A;Molecule type: DNA  
A;Residues: 1-301 <JOS>  
A;Cross-references: EMBL:X60432; NID:G433706; PIDN:CAA2959.1; PID:G433707  
C;Superfamily: hydroxyproline-rich glycoprotein  
C;Experimental source: strain W64A

Query Match	60.6%;	Score 54.5;	DB 2;	Length 301;
Best Local Similarity	71.4%;	Pred. No. 9.7;		
Matches 10;	Conservative	1;	Mismatches	2;
Matches 1;	Indels	1;	Gaps	1;

Qy 3 RPRPYL-PRPRPP 15  
|||:||||  
Db 149 RPSPPVPTPRPP 16

## RESULT 9

T113478  
hypothetical protein 34P3.10 - fruit fly (*Drosophila melanogaster*)  
C:Species: *Drosophila melanogaster*  
C:date: 13-Aug-1999 #sequence\_revision 13-Aug-1999 #text\_change 17-Nov-2000  
C:Accession: T113478  
R:Valenti, P.; Salles, C.; Campbell, L.; Glover, D.  
Submitted to the EMBL Data Library, April 1999  
A:Description: Sequencing the distal X chromosome of *Drosophila melanogaster*.  
A:Reference number: Z17685

A;Accession: U5478  
A;Status: preliminary; translated from GB/EMBL/DBJ  
A;Molecule type: DNA

A;Molecule type: DNA  
A;Residues: 1-359 <PH>  
A;Cross-references: EMBL:AL031583; NID:e1321005; PID:e1321018; PIDN:CAB41346.1  
C;Genetics:  
A;Cross-references: FlyBase:FBgn0025623  
A;Introns: 17/2; 50/3; 333/2  
A;Note: EG:34F3.10

Query Match	60.0%;	Score 54;	DB 2;	Length 359;
Best Local Similarity	71.4%;	Pred. NO. 13;		
Matches 10;	Conservative	0;	Mismatches	4;
			Indels	0;
			Gaps	0;

Qy 2 RRPRPPYLRPRPP 15  
||| ||| ||| |||  
Db 167 RRPPPPPLPPPPPP 180

## RESULT 10

T32652  
hypothetical protein F39C12.3 - *Caenorhabditis elegans*  
C;Species: *Caenorhabditis elegans*  
C;Date: 29-Oct-1999 #sequence\_revision 29-Oct-1999 #text\_change 18-Feb-2000  
C;Accession: T32652  
R;Chissoe, S.; Sansone, J.  
submitted to the EMBL Data Library, December 1997  
A;Description: The sequence of *C. elegans* cosmid F39C12.  
A;Reference number: Z1206  
A;Accession: T32652  
A;Status: preliminary;  
A;Molecule type: DNA  
translated from GB/EMBL/DBDJ

A;Residues: 1-427 <CHI>  
 A;Cross-references: EMBL:AF039043; PIDN:AAB94196.1; GSPDB:GN00028; CESP:F39C12.3  
 A;Experimental source: strain Bristol N2; clone F39C12  
 C;Genetics:  
 A;Gene: CESP:F39C12.3  
 A;Map position: X  
 A;Introns: 42/3; 104/3; 133/3; 164/3; 213/3; 276/3; 336/3

Query Match 60.0%; Score 54; DB 2; Length 427;  
 Best Local Similarity 69.2%; Pred. No. 16;

Matches 9; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRPVLPRLPRPP 15

|||||:|||||

Db 338 RRPDPDIPPLPP 350

RESULT 11

S06675

apidaecin Ib precursor - honeybee

C;Species: Apis mellifera (honeybee)

C;Date: 07-Sep-1990 #sequence\_revision 07-Sep-1990 #text\_change 16-Dec-1998

C;Accession: S06675

R;Casteels-Josson, K.; Capaci, T.; Tempst, P.

EMBO J. 8, 2387-2391, 1989

A;Title: Apidaecins: antibacterial peptides from honeybees.

A;Reference number: S05383; MUID:90005446; PMID:2676519

A;Accession: S06675

A;Molecule type: protein

A;Residues: 1-26 <CAS>

F;1-8/Domain: propeptide #status experimental <PRO>

F;9-26/Product: apidaecin Ib #status experimental <MAT>

Query Match 58.9%; Score 53; DB 2; Length 26;

Best Local Similarity 72.7%; Pred. No. 1.3;

Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLPRLPRPP 15

|||||:|||||

Db 12 RPYVLPRLPRPP 22

RESULT 12

S35331

apidaecin 22 precursor - honeybee

C;Species: Apis mellifera (honeybee)

C;Date: 03-Feb-1994 #sequence\_revision 03-Feb-1994 #text\_change 21-Jul-2000

C;Accession: S35331

R;Casteels-Josson, K.; Capaci, T.; Tempst, P.

EMBO J. 12, 1569-1578, 1993

A;Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplification

A;Reference number: S35330; MUID:93223697; PMID:8467807

A;Accession: S35331

A;Status: preliminary

A;Molecule type: DNA

A;Residues: 1-144 <CAS>

A;Cross-references: EMBL:X72576; NID:9297064; PIDN:CAA51168.1; PID:9297065

C;Superfamily: procyclic acidic repetitive protein

Query Match 58.9%; Score 53; DB 2; Length 144;

Best Local Similarity 72.7%; Pred. No. 7.1;

Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLPRLPRPP 15

|||||:|||||

Db 102 RPYVLPRLPRPP 112

RESULT 13

T29373

hypothetical protein ZC404.1 - Caenorhabditis elegans

C;Species: Caenorhabditis elegans

C;Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 09-Jun-2000

C;Accession: T29373  
 R;Bentley, D.; Le, T.T.  
 submitted to the EMBL Data Library, April 1996  
 A;Description: The sequence of C. elegans cosmid ZC404.  
 A;Reference number: Z20614  
 A;Accession: T29373  
 A;Status: preliminary; translated from GB/EMBL/DBJ  
 A;Molecule type: DNA  
 A;Residues: 1-184 <BEN>  
 A;Cross-references: EMBL:U55363; PIDN:AAA97967.1; GSPDB:GN00023; CESP:ZC404.1  
 A;Experimental source: strain Bristol N2; clone ZC404  
 C;Genetics:  
 A;Gene: CESP:ZC404.1  
 A;Map position: 5  
 A;Introns: 15/2; 50/2; 75/2; 138/2  
 C;Superfamily: Caenorhabditis elegans hypothetical protein ZC404.1

Query Match 58.9%; Score 53; DB 2; Length 184;

Best Local Similarity 90.0%; Pred. No. 9;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 RRPVLPRLPR 12

|||||:|||||

Db 26 RPKKPYLP 35

RESULT 14

S35332

apidaecin 73 precursor - honeybee (fragment)

N;Contains: apidaecin Ia

C;Species: Apis mellifera (honeybee)

C;Date: 03-Feb-1994 #sequence\_revision 03-Feb-1994 #text\_change 03-Nov-2000

C;Accession: S35332; S05383

R;Casteels-Josson, K.; Capaci, T.; Tempst, P.

EMBO J. 12, 1569-1578, 1993

A;Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplification

A;Reference number: S35330; MUID:93223697; PMID:8467807

A;Accession: S35332

A;Molecule type: mRNA

A;Residues: 1-283 <CAS>

A;Cross-references: EMBL:X72577; NID:9297066; PIDN:CAA51169.1; PID:94539289

A;Accession: S05383

A;Molecule type: protein

A;Residues: 258-283 <CA3>

C;Superfamily: proline-rich protein

F;266-283/Product: apidaecin Ia #status experimental <MAT>

Query Match 58.9%; Score 53; DB 2; Length 283;

Best Local Similarity 72.7%; Pred. No. 14;

Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 5 RPYVLPRLPRPP 15

|||||:|||||

Db 241 RPYVLPRLPRPP 251

RESULT 15

E71415

probable coll wall protein - Arabidopsis thaliana

C;Species: Arabidopsis thaliana (mouse-ear cress)

A;Variety: Columbia

C;Date: 03-Aug-1998 #sequence\_revision 03-Aug-1998 #text\_change 05-Dec-1998

C;Accession: E71415

R;Bevan, M.; Bancroft, I.; Bent, E.; Love, K.; Goodman, H.; Dean, C.; Bergkamp, R.; Dirks

P.; Wedler, H.; Wedler, E.; Wambutt, T.; Weitzenecker, T.; Pohl, T.M.; Terry, N.; Giele

avanagh, T.; Hempel, S.; Kotter, P.; Entian, K.D.; Rieger, M.; Schaeffer, M.; Funk, B.

Nature 391, 485-488, 1998

A;Authors: Mueller-Auer, S.; Silvey, M.; James, R.; Montfort, A.; Pons, A.; Puigdomenech,

erhoft, A.; Moores, T.; Jones, J.D.G.; Eneva, T.; Palme, K.; Benes, V.; Rechman, S.; Ansc

C.; Chalwatzis, N.

A;Title: Analysis of 1.9 Mb of contiguous sequence from chromosome 4 of Arabidopsis thali

A;Reference number: A71400; MUID:98121113; PMID:9461215

A;Accession: E71415

A;Status: preliminary; nucleic acid sequence not shown; translation not shown  
A;Molecule type: DNA  
A;Residues: 1-428 <BEV>  
A;Cross-references: GB:Z97338; NID:G2244870; PID:G327461; PID:G224487#  
C;Genetics:  
A;Map position: 4COP9-4G3845

Query Match 58.9%; Score 53; DB 2; Length 428;  
Best Local Similarity 61.5%; Pred. No. 21;  
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;  
Qy 3 RPRPPYLP RPRPP 15  
Db 67 KPPPPYIPCP RPP 79

Search completed: May 13, 2003, 10:41:48  
Job time : 18 secs

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GenCore version 5.1.4 p5 4578  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: May 13, 2003, 10:34:02 ; Search time 11 Seconds  
(without alignments)  
.56.559 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPRPPYLPRPP 15

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt\_40.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	90	100.0	172	1 PR39_PIG	P80054 sus scrofa
2	66	73.3	190	1 BCT7_BOVIN	P19661 bos taurus
3	59.5	66.1	107	1 COTT_BACSU	P11863 bacillus su
4	58	64.4	190	1 BCT7_SHEEP	P50415 ovis aries
5	56.5	62.8	168	1 AP14_APIME	Q06601 apis mellif
6	55.5	61.7	151	1 RNB_HSV2H	P89479 herpes simp
7	53	58.9	144	1 AP22_APIME	P35581 apis mellif
8	53	58.9	283	1 AP73_APIME	Q06602 apis mellif
9	53	58.9	381	1 PRLP_BOVIN	Q9gk88 bos taurus
10	52	57.8	261	1 RLI_HSV2H	P28283 herpes simp
11	52	57.8	354	1 ATHI_HUMAN	Q92858 homo sapien
12	52	57.8	467	1 AFCE_ARATH	P51566 arabidopsis
13	52	57.8	841	1 RELA_STRAT	Q85709 streptomyce
14	51.5	57.2	1187	1 PTNE_HUMAN	Q15678 homo sapien
15	51.5	57.2	1189	1 PTNE_MOUSE	Q62130 mus musculu
16	51	56.7	15	1 MK1_PALPR	P80408 palomela pr
17	51	56.7	180	1 XG_HUMAN	P55808 homo sapien
18	50.5	56.1	393	1 C1W4_HUMAN	Q9ny98 homo sapien
19	50	55.6	17	1 APID_BOMPA	P81464 bombus pasc
20	50	55.6	415	1 ACRO_PIG	P08001 sus scrofa
21	50	55.6	424	1 S3B4_HUMAN	Q15427 homo sapien
22	50	55.6	449	1 APG_BRANA	P40603 brassica na
23	50	55.6	678	1 ABPE_RIPCL	Q27905 riptortus c
24	49	54.4	134	1 PRL5_HUMAN	Q99954 homo sapien
25	49	54.4	296	1 GDA6_WHEAT	P04726 triticum ae
26	49	54.4	352	1 RRS1_ARATH	Q9sh88 arabidopsis
27	49	54.4	2911	1 FBN2_HUMAN	P35556 homo sapien
28	48.5	53.9	2142	1 BAT2_HUMAN	P48634 homo sapien
29	48	53.3	280	1 TNF6_CERTO	Q9bdn1 cercocebus
30	48	53.3	280	1 TNF6_MACMU	Q9myl6 macaca mula
31	48	53.3	281	1 TNF6_HUMAN	P48023 homo sapien
32	48	53.3	282	1 TNF6_PIG	Q9bea8 sus scrofa
33	48	53.3	402	1 VGLD_PRVRI	P07645 pseudorabie

#### RESULT 1

ID	PR39_PIG	STANDARD;	PRT;	172 AA.
AC	P80054; Q9TR84;			
DT	01-MAR-1992 (Rel. 21, Created)			
DT	01-OCT-1996 (Rel. 34, Last sequence update)			
DT	16-OCT-2001 (Rel. 40, Last annotation update)			
DE	Antibacterial protein PR-39 precursor.			
GN	PR39.			
OS	Sus scrofa (Pig).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.			
OX	NCBI_TaxID=9823;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=95350216; PubMed=7624374;			
RA	Gudmundason G.H., Magnusson K.P., Chowdhary B.P., Johansson M., Andersson L., Boman H.G.;			
RT	"Structure of the gene for porcine peptide antibiotic PR-39, a cathelin gene family member: comparative mapping of the locus for the human peptide antibiotic FALL-39."			
RT	Proc. Natl. Acad. Sci. U.S.A. 92:7085-7089(1995).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Bone marrow;			
RX	MEDLINE=94071853; PubMed=8250863;			
RA	Storici P., Zanetti M.;			
RT	"A cDNA derived from pig bone marrow cells predicts a sequence identical to the intestinal antibacterial peptide PR-39."			
RL	Biochem. Biophys. Res. Commun. 196:1058-1065(1993).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Liver;			
RX	MEDLINE=96105365; PubMed=7498526;			
RA	Zhao C., Ganz T., Lehrer R.I.;			
RT	"Structures of genes for two cathelin-associated antimicrobial peptides: prophenin-2 and PR-39."			
RL	FEBS Lett. 376:130-134(1995).			
RN	[4]			
RP	SEQUENCE OF 131-169.			
RC	TISSUE=Intestine;			
RX	MEDLINE=92111534; PubMed=1765098;			
RA	Agerberth B., Lee J.-Y., Bergman T., Carlquist M., Boman H.G., Mutt V., Joernvall H.;			
RT	"Amino acid sequence of PR-39. Isolation from pig intestine of a new member of the family of proline-arginine-rich antibacterial peptides."			
RL	Eur. J. Biochem. 202:849-854(1991).			
RN	[5]			
RP	SEQUENCE OF 131-164, AND FUNCTION.			
RC	TISSUE=Neutrophils;			
RX	MEDLINE=95088504; PubMed=7996056;			
RA	Shi J., Ross C.R., Chengappa M.M., Blecha F.;			
RT	"Identification of a proline-arginine-rich antibacterial peptide from neutrophils that is analogous to PR-39, an antibacterial peptide from the small intestine."			

#### ALIGNMENTS

34	48	53.3	520	1 C84A_ARATH	Q42600 arabidopsis
35	48	53.3	759	1 TOP3_CAEEL	O61660 caenorhabdi
36	47.5	52.8	283	1 EXTN_SORBI	P24152 sorghum bic
37	47.5	52.8	372	1 DBPA_HUMAN	P16989 homo sapien
38	47.5	52.8	1443	1 SYO2_HUMAN	O15056 homo sapien
39	47	52.2	176	1 BCT5_BOVIN	P19660 bos taurus
40	47	52.2	261	1 PRP2_MOUSE	P05142 mus musculu
41	47	52.2	296	1 PRP3_MOUSE	P05143 mus musculu
42	47	52.2	507	1 MEFA_HUMAN	Q02078 homo sapien
43	47	52.2	753	1 SX30_HUMAN	O94993 homo sapien
44	47	52.2	846	1 IRS1_HCMVA	P09715 human cytom
45	47	52.2	1040	1 BOI2_YEAST	P39969 saccharomyc





```

OX NCBI_TaxID=1423;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=168 / JH642;
RX MEDLINE=89313296; PubMed=2546006;
RA Aronson A.I., Song H.Y., Bourne N.;
RT "Gene structure and precursor processing of a novel Bacillus subtilis
  spore coat protein.";
RL Mol. Microbiol. 3:437-444 (1989).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=168;
RX MEDLINE=98044033; PubMed=9384377;
RA Kunst F., Ogasawara N., Mosser I., Albertini A.M., Alloni G.,
  Azevedo V., Bertetto M.G., Bessieres P., Bolotin A., Borchert S.,
  Borriass R., Boursier L., Brans A., Braun M., Brignell S.C., Bron S.,
  Brouillet S., Bruschini C.V., Caldwell B., Capuano V., Carter N.M.,
  Choi S.K., Codani J.J., Conneron I.F., Cummings N.J., Daniel R.A.,
  Denizot F., Devine K.M., Dusterhoft A., Ehrlich S.D., Emmerson P.T.,
  Entian K.D., Errington J., Fabret C., Ferrari E., Foulger D.,
  Fritz C., Fujita M., Fujita Y., Fuma S., Galizzi A., Galleron N.,
  Chim S.Y., Glaser P., Goffeau A., Gollightly E.J., Grandi G.,
  Guisepi G., Gay B.J., Haga K., Hatach J., Harwood C.R., Henaut A.,
  Hilbert H., Holsappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,
  Joris B., Karamata D., Kasahara Y., Klaerr-Blanchard M., Klein C.,
  Kobayashi Y., Koetter P., Koningstein G., Krogh S., Kumano M.,
  Kurita K., Lapidus A., Lardinois S., Lauber J., Lazarevic V.,
  Lee S.M., Levine A., Liu H., Masuda S., Mael C., Medigue C.,
  Medina N., Mellado R.P., Mizuno M., Moestl D., Nakai S., Noback M.,
  Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,
  Parro V., Pohl T.M., Portetelle D., Porwollik S., Prescott A.M.,
  Pressac E., Pujic P., Purnelle B., Rapoport G., Rey M., Reynolds S.,
  Rieger M., Rivolta C., Rocha E., Roche B., Rose M., Sadaie Y.,
  Sato T., Scanlan E., Schleich S., Schroeter R., Scoffone F.,
  Sekiguchi J., Sekowska A., Seror S.J., Serron P., Shin B.S., Soldo B.,
  Sorokin A., Tacconi E., Takagi T., Takahashi H., Takemaru K.,
  Takeuchi M., Tamakoshi A., Tanaka T., Terpstra P., Tognoni A.,
  Tosato V., Uchiyama S., Vandenbol M., Vannier F., Vassarotti A.,
  Viari A., Wambitt R., Wedler E., Wedler H., Weitzensger T.,
  Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
  Yoshida K., Yoshikawa H.F., Zumbstein E., Yoshikawa H., Danchin A.;
RT "The complete genome sequence of the Gram-positive bacterium Bacillus
  subtilis.";
RL Nature 390:249-256 (1997).
CC -!- FUNCTION: POSSIBLY PROTECTION OF SPORE AND PROBABLY PLAYS
  SOME ROLE IN GERMINATION.
CC -!- SUBCELLULAR LOCATION: OUTER SURFACE OF ENDOSPORE.
CC
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  or send an email to license@isb-sib.ch).
CC
DR EMBL; X13740; CAA32004.1; -
DR EMBL; Z99110; CAB32066.1; -
DR FIR; S04835; S04835.
DR Subtilist; BG10495; cotT.
KW Sporulation; Signal; Complete proteome.
FT SIGNAL 1 44
FT CHAIN 45 107 SPORE COAT PROTEIN T.
SQ SEQUENCE 107 AA; 12992 MW; AD1P66F0C4CE29A3 CRC64;

Query Match 66.1%; Score 59.5; DB 1; Length 107;
Best Local Similarity 84.6%; Pred. No. 0.47;
Matches 11; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 4 PRPP-YLPRPRPP 15
DB 74 PRPPYYPRPRPP 86

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RESULT 4
BCT7_SHEEP
ID BCT7_SHEEP STANDARD; PRT; 190 AA.
AC P50415;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE Bactenein 7 precursor (BAC7).
GN BAC7.5.
OS Ovis aries (Sheep).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Caprinae; Ovis.
OX NCBI_TaxID=9940;
RN [1]_TaxID=9940;
RP SEQUENCE FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=96105386; PubMed=7498547;
RA Bagella L., Scocchi M., Zanetti M.;
RT "cDNA sequences of three sheep myeloid cathelicidins.";
RL FEBS Lett. 376:225-228 (1995).
CC -!- FUNCTION: EXERTS, IN VITRO, A POTENT ANTIMICROBIAL ACTIVITY.
  PROBABLY DUE TO AN IMPAIRMENT OF THE FUNCTION OF THE RESPIRATORY
  CHAIN AND OF ENERGY-DEPENDENT ACTIVITIES IN THE INNER MEMBRANE
  OF SUSCEPTIBLE MICROORGANISMS (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.
CC
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CC
DR EMBL; L46852; AAA85468.1; -
DR InterPro; IPR001894; Cathelicidin.
DR Pfam; PF00666; Cathelicidins; 1.
DR ProDom; PD001838; Cathelicidin; 1.
DR PROSITE; PS00946; CATHELICIDINS_1; 1.
DR PROSITE; PS00947; CATHELICIDINS_2; 1.
KW Antibiotic; Repeat; Signal.
FT SIGNAL 1 29 POTENTIAL.
FT PROPEP 30 130 BY SIMILARITY.
FT CHAIN 131 190 BACTENECIN 7.
FT MOD_RES 30 30 PYRROLIDONE CARBOXYLIC ACID
  (BY SIMILARITY).
FT DISULFID 85 96 BY SIMILARITY.
FT DISULFID 107 124 BY SIMILARITY.
SQ SEQUENCE 190 AA; 21829 MW; E4AAFB1600E98371 CRC64;

Query Match 64.4%; Score 58; DB 1; Length 190;
Best Local Similarity 78.6%; Pred. No. 1.2;
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RRRPRPPYLRPRPP 14
DB 132 RLRRPRRLPRPRPP 145

RESULT 5
API4_APIME
ID API4_APIME STANDARD; PRT; 168 AA.
AC Q06601; P11525; P11526; P11527;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE Apidaecin precursor, type 14.
GN API4.
OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
OC Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;

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OC ACuleata; Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;
RN SEQUENCE FROM N.A.
RP MEDLINE=93223697; PubMed=8467807;
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RX "Apidaecin multipetide precursor structure: a putative mechanism for
RT amplification of the insect antibacterial response.";
RL EMBO J. 12:1569-1578(1993).
RN [2]
RP SEQUENCE OF APIDAECINS IA/IB/II.
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMBO J. 8:2387-2391(1989).
CC -!- FUNCTION: APIDAECINS HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY
CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
CC PROPAGATION.
CC
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CC
CC EMBL: X72575; CAA51167.1; -.
DR PIR: S05383; S05383.
DR PIR: S06675; S06675.
DR PIR: S06676; S06676.
DR PIR: S35330; S35330.
DR InterPro: IPR004828; Apidaecin.
DR Pfam: PF00807; Apidaecin; 5.
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
KW Cleavage on pair of basic residues; Repeat.
FT SIGNAL 1 19
FT PROPEP 35 42
FT PEPTIDE 43 60
FT PROPEP 63 70
FT PEPTIDE 71 88
FT PROPEP 91 98
FT PEPTIDE 99 116
FT PROPEP 119 124
FT PEPTIDE 125 142
FT PROPEP 145 150
FT PEPTIDE 151 168
SQ SEQUENCE 168 AA; 19380 MW; 594B931254C04A37 CRC64;
Query Match 62.8%; Score 56.5; DB 1; Length 168;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 11; Conservative 2; Mismatches 2; Indels 7; Gaps 1;
OY 1 RRRP-----RPPYLPRLPRPP 15
Db 117 RREPEAEFGNNRPVYIQPRPP 138
RESULT 6
RNB_HSV2H ID_RNB_HSV2H STANDARD; PRT; 151 AA.
AC P9479;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Potential RNA-binding protein.
GN US11.
OS Herpes simplex virus (type 2 / strain HG52).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Simplexvirus.
OX NCBI_TaxID=10315;
RN [1]
RP SEQUENCE FROM N.A.
RA Dolan A.;
RL Submitted (FEB-1997) to the EMBL/GenBank/DBSJ databases.
CC -!- FUNCTION: BINDS DNA AND RNA (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: Nuclear; nucleolar (By similarity).
CC
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CC
CC EMBL: Z86099; CAB06719.1; -.
DR DNA-binding; RNA-binding; Repeat; Nuclear protein.
FT DOMAIN 90 146
FT REPEAT 90 95
FT REPEAT 96 101
FT REPEAT 102 104
FT REPEAT 105 110
FT REPEAT 111 116
FT REPEAT 117 122
FT REPEAT 123 128
FT REPEAT 129 130
FT REPEAT 131 134
FT REPEAT 135 140
FT REPEAT 141 146
SQ SEQUENCE 151 AA; 16297 MW; FAB751F23C3DB6AE CRC64;
Query Match 61.7%; Score 55.5; DB 1; Length 151;
Best Local Similarity 73.3%; Pred. No. 1.9;
Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;
OY 2 RRPRLPRLPR-PRPP 15
Db 127 RPPRLPRLPRLPRPP 141
RESULT 7
AP22_APIME ID AP22_APIME STANDARD; PRT; 144 AA.
AC P35581; P11525; P11526;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE Apidaecin precursor, type 22.
OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;
OC Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;
OC Aculeata; Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;
RN SEQUENCE FROM N.A.
RP MEDLINE=93223697; PubMed=8467807;
RX Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RA "Apidaecin multipetide precursor structure: a putative mechanism for
RT amplification of the insect antibacterial response.";
RL EMBO J. 12:1569-1578(1993).
RN [2]
RP SEQUENCE (APIDAECINS IA/IB).
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMBO J. 8:2387-2391(1989).
CC -!- FUNCTION: APIDAECINS HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY
CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
CC PROPAGATION.
CC
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DR EMBL; X72576; CAA51168.1; -.  
DR PIR; S05383; S05383.  
DR PIR; S06675; S06675.  
DR PIR; S35331; S35331.  
DR InterPro: IPR004828; Apidaecin.  
DR Pfam: PF00807; Apidaecin 4.  
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;  
KW Cleavage on pair of basic residues; Repeat.  
FT SIGNAL 1 19 POTENTIAL.  
FT PROPEP 35 42  
FT PEPTIDE 43 60 APIDAEACIN IB.  
FT PROPEP 63 70  
FT PEPTIDE 71 88 APIDAEACIN IB.  
FT PROPEP 91 98  
FT PEPTIDE 99 116 APIDAEACIN IB.  
FT PROPEP 119 126  
FT PEPTIDE 127 144 APIDAEACIN IA.  
SQ SEQUENCE 144 AA; 16539 MW; 6FA1AD74CB77108D CRC64;

Query Match 58.9%; Score 53; DB 1; Length 144;  
Best Local Similarity 72.7%; Pred. No. 3.6;  
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 5 RPPYLP RP RP 15

DB 102 RPYIIPQRP 112

#### RESULT 8

AP73 APIME STANDARD; PRT; 283 AA.  
AC Q06602; P11525; P11526;  
DT 01-OCT-1989 (Rel. 12, Created)  
DT 01-JUN-1994 (Rel. 29, Last sequence update)  
DT 01-JUN-1994 (Rel. 29, Last annotation update)  
DE Apidaecin precursor, type 73 (Fragment).  
GN APID73.  
OS Apis mellifera (Honeybee).  
OC Eukaryota; Metazoa; Arthropoda; Mandibulata; Pancrustacea; Hexapoda;  
OC Insecta; Pserygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;  
OC Aculeata; Apoidea; Apidae; Apis.  
OX NCBI\_TaxID=7460;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=93223697; PubMed=8467807;  
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;  
RT "Apidaecin multipetide precursor structure: a putative mechanism for  
RT amplification of the insect antibacterial response.";  
RL EMBO J. 12:1569-1578(1993).  
RN [2]  
RP SEQUENCE OF APIDAEACIN IA/IB.  
RC TISSUE=Hemolymph; PubMed=2676519;  
RX MEDLINE=90005446; PubMed=2676519;  
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;  
RT "Apidaecins: antibacterial peptides from honeybees.";  
RL EMBO J. 8:2387-2391(1989).  
CC -!- FUNCTION: APIDAEACIN HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY  
CC AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL  
CC PROPAGATION.  
CC -----

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CC -----

DR EMBL; X72577; CAA51169.1; -.  
DR PIR; S05383; S05383.  
DR PIR; S06675; S06675.  
DR PIR; S35332; S35332.  
DR InterPro: IPR004828; Apidaecin.  
DR Pfam: PF00807; Apidaecin 9.  
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;  
KW Cleavage on pair of basic residues; Repeat.  
FT NON\_TER 1 18 POTENTIAL.  
FT SIGNAL <1 18  
FT PROPEP 34 41 APIDAEACIN IB.  
FT PEPTIDE 42 59  
FT PROPEP 62 69  
FT PEPTIDE 70 87 APIDAEACIN IB.  
FT PROPEP 90 97  
FT PEPTIDE 98 115 APIDAEACIN.  
FT PROPEP 118 125  
FT PEPTIDE 126 143 APIDAEACIN IB.  
FT PROPEP 146 153  
FT PEPTIDE 154 171 APIDAEACIN.  
FT PROPEP 174 182  
FT PEPTIDE 183 199 APIDAEACIN IB.  
FT PROPEP 202 209  
FT PEPTIDE 210 227 APIDAEACIN IB.  
FT PROPEP 230 237  
FT PEPTIDE 238 255 APIDAEACIN IB.  
FT PROPEP 258 265  
FT PEPTIDE 266 283 APIDAEACIN IA.  
SQ SEQUENCE 283 AA; 32695 MW; 4EA5FDECD5E142B CRC64;

Query Match 58.9%; Score 53; DB 1; Length 283;  
Best Local Similarity 72.7%; Pred. No. 6.9;  
Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 5 RPPYLP RP RP 15

DB 241 RPYIIPQRP 251

#### RESULT 9

PRLP BOVIN STANDARD; PRT; 381 AA.  
ID PRLP BOVIN  
AC Q9GKN8;  
DT 15-JUN-2002 (Rel. 41, Created)  
DT 15-JUN-2002 (Rel. 41, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE Proargin precursor (Proline-arginine-rich end leucine-rich repeat  
DE of protein).  
GN PRELP.  
OS Bos taurus (Bovine).  
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
OC Bovidae; Bovinae; Bos.  
OX NCBI\_TaxID=9913;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC TISSUE=Articular cartilage;  
RX MEDLINE=20576219; PubMed=11007795;  
RA Bengtsson E., Asberg A., Heinegaard D., Sommarin Y., Spillmann D.;  
RT "The amino-terminal part of PRELP binds to heparin and heparan  
RT sulfate.";  
RL J. Biol. Chem. 275:40695-40702(2000).  
RN [2]  
RP FUNCTION.  
RX PubMed=11847210;  
RA Bengtsson E., Moergelin M., Sasaki T., Timpl R., Heinegaard D.,  
RA Asberg A.;  
RT "The leucine-rich repeat protein PRELP binds perlecan and collagens  
RT and may function as a basement membrane anchor.";  
RL J. Biol. Chem. 277:15061-15068(2002).  
CC -!- FUNCTION: May anchor basement membranes to the underlying  
CC connective tissue.  
CC -!- SUBUNIT: Binds the basement membrane heparan sulfate proteoglycan

perlecan and triple helical collagens type I and type II.  
-1- SUBCELLULAR LOCATION: Extracellular matrix.  
-1- DOMAIN: The basic amino-terminal Arg/Pro-rich binds heparin and heparan sulfate. Binds collagens type I and type II through its leucine-rich repeat domain.  
-1- SIMILARITY: BELONGS TO THE SMALL LEUCINE-RICH PROTEOGLYCANS (SLRPs) FAMILY, CLASS II SUBFAMILY.  
-1- SIMILARITY: CONTAINS 12 LEUCINE-RICH REPEATS (LRR).  
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EMBL: AF163568; AAG23723.1; -  
InterPro: IPR001611; LRR.  
InterPro: IPR003372; LRR Nterm.  
InterPro: IPR003592; LRR C-  
InterPro: IPR003591; LRR C-  
Pfam: PF00560; LRR; 10.  
Pfam: PF01462; LRRNT; 1.  
PRINTS: PR00019; LEURICHRPT.  
SMART: SM00370; LRR; 7.  
SMART: SM00013; LRRNT; 1.  
SMART: SM00369; LRR-TIP; 7.  
KW Glycoprotein; Extracellular matrix; Repeat; Leucine-rich repeat; Signal.  
KW SIGNAL.  
FT CHAIN 1 21 POTENTIAL.  
FT CHAIN 22 381 PROLARGIN.  
FT DOMAIN 72 88 CYS-RICH.  
FT REPEAT 94 113 LRR-S 1.  
FT REPEAT 114 137 LRR-T 1.  
FT REPEAT 138 161 LRR-T 2.  
FT REPEAT 162 182 LRR-S 2.  
FT REPEAT 183 206 LRR-T 3.  
FT REPEAT 207 232 LRR-T 4.  
FT REPEAT 233 253 LRR-S 3.  
FT REPEAT 254 277 LRR-T 5.  
FT REPEAT 278 302 LRR-T 6.  
FT REPEAT 303 322 LRR-S 4.  
FT REPEAT 323 361 LRR-T 7.  
FT REPEAT 362 381 LRR-T 8.  
FT DOMAIN 196 201 POLY-LEU.  
FT DISULFID 331 372 BY SIMILARITY.  
FT CARBOHYD 123 123 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 288 288 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 319 319 N-LINKED (GLCNAC. . .) (POTENTIAL).  
FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).  
SQ SEQUENCE 381 AA; 43682 MW; 23DA99C01B772A0 CRC64;  
-----  
Query Match 58.9%; Score 53; DB 1; Length 381;  
Best Local Similarity 76.9%; Pred. No. 9.2;  
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
QY 2 RRRPPVLPVRPP 14  
DB 25 RRRPPVLPVRPP 37  
-----  
RESULT 10  
RLA\_HSV2H STANDARD; PRT; 261 AA.  
AC P28283;  
DT 01-DEC-1992 (Rel. 24, Created)  
DT 01-DEC-1992 (Rel. 24, Last sequence update)  
DT 01-NOV-1997 (Rel. 35, Last annotation update)  
DE Neurovirence factor (ICP34.5).  
GN RL1.  
OS Herpes simplex virus (type 2 / strain HG52).  
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;

Alphaherpesvirinae; Simplexvirus.  
NCBI\_TaxID=10315;  
[1]  
SEQUENCE FROM N.A.  
MEDLINE=92113549; PubMed=1662697;  
RA McGeech D.J., Cunningham C., McIntyre G., Dolan A.;  
RT "Comparative sequence analysis of the long repeat regions and adjoining parts of the long unique regions in the genomes of herpes simplex viruses types 1 and 2";  
RT J. Gen. Virol. 72:3057-3075(1991).  
[2]  
SEQUENCE FROM N.A.  
RA Dolan A.;  
RL Submitted (FEB-1997) to the EMBL/GenBank/DBJ databases.  
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EMBL: D10471; BAA23428.1; -  
DR EMBL; 286099; CAB06759.1; -  
DR EMBL; 286099; CAB06706.1; -  
DR PIR; JQ1502; WMBEXE.  
KW Repeat.  
FT DOMAIN 3 12 2 X 5 AA TANDEM REPEATS OF R-R-R-G-P.  
FT REPEAT 3 7  
FT REPEAT 8 12 2 X 8 AA TANDEM REPEATS OF P-R-P-G-A-P-A-  
FT DOMAIN 16 31 V.  
FT REPEAT 16 23  
FT REPEAT 24 31  
SQ SEQUENCE 261 AA; 27908 MW; 4BBD13AF3D906D71 CRC64;  
-----  
Query Match 57.8%; Score 52; DB 1; Length 261;  
Best Local Similarity 64.7%; Pred. No. 8.3;  
Matches 11; Conservative 1; Mismatches 3; Indels 2; Gaps 1;  
QY 1 RRRPRP--PYLPRPRPP 15  
DB 13 RRRPRPGAPVPRPGAP 29  
-----  
RESULT 11  
ATH1\_HUMAN STANDARD; PRT; 354 AA.  
AC Q92858;  
DT 15-DEC-1998 (Rel. 37, Created)  
DT 15-DEC-1998 (Rel. 37, Last sequence update)  
DT 15-DEC-1998 (Rel. 37, Last annotation update)  
DE Atonal protein homolog 1 (Helix-loop-helix protein hATH-1).  
GN ATOH1 OR ATH1.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
OC NCBI\_TaxID=9606;  
[1]  
SEQUENCE FROM N.A.  
RA Ben-Arie N., McCall A.E., Berkman S., Eichele G., Bellen H.J., Zoghbi H.Y.;  
RT "Evolutionary conservation of sequence and expression of the bHLH protein Atonal suggests a conserved role in neurogenesis.";  
RL Hum. Mol. Genet. 5:1207-1216(1996).  
CC -1- FUNCTION: ACTIVATES E BOX-DEPENDENT TRANSCRIPTION IN COLLABORATION WITH E47, BUT THE ACTIVITY IS COMPLETELY ANTAGONIZED BY THE NEGATIVE REGULATOR OF NEUROGENESIS HES-1. MAY PLAY A ROLE IN THE DIFFERENTIATION OF SUBSETS OF NEURAL CELLS BY ACTIVATING E BOX-DEPENDENT TRANSCRIPTION (BY SIMILARITY).  
CC -1- SUBUNIT: EFFICIENT DNA BINDING REQUIRES DIMERIZATION WITH ANOTHER

```

CC BHLH PROTEIN.
CC -!- SUBCELLULAR LOCATION: Nuclear (Probable).
CC -!- SIMILARITY: BELONGS TO THE BASIC HELIX-LOOP-HELIX (BHLH) FAMILY OF
CC TRANSCRIPTION FACTORS. "ATONAL" SUBFAMILY.
CC -----
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CC -----
CC EMBL; U61148; AAB41305.1; -.
CC DR TRANSFAC; T04544; -.
CC DR Genew; HGNC:797; ATOH1.
CC DR MIM; 601461; -.
CC DR InterPro; IPR001092; HLH_basic.
CC DR Pfam; PF00010; HLH; 1.
CC DR SMART; SM00353; HLH; 1.
CC DR PROSITE; PS00038; HLH_1; FALSE_NEG.
CC DR PROSITE; PS0888; HLH_2; 1.
CC KW Transcription regulation; Activator; DNA-binding; Nuclear protein.
CC FT DOMAIN 29 38 POLY-PRO.
CC FT DNA_BIND 160 171 BASIC DOMAIN.
CC FT DOMAIN 172 212 HELIX-LOOP-HELIX MOTIF (BY SIMILARITY).
CC FT DOMAIN 224 228 POLY-PRO.
CC SQ SEQUENCE 354 AA; 39160 MW; AB12F1E917A00A8D CRC64;

Query Match 57.8%; Score 52; DB 1; Length 354;
Best Local Similarity 57.1%; Pred. NO. 11;
Matches 8; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RRRPPVLPRLPP 15
|:|:|:|:|:|
DB 21 *RQPHLPQPPPP 34

RESULT 12
AFCL_ARATH
ID AFCL_ARATH STANDARD; PRT; 467 AA.
AC P51566;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Protein kinase AFCL (EC 2.7.1.1.-).
GN AFCL.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopses.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Landsberg erecta;
RX MEDLINE=95083650; PubMed=7991592;
RA Bender J., Fink G.R.;
RT "AFCL, a LAMMER kinase from Arabidopsis thaliana, activates STE12-
RT dependent processes in yeast.";
RL Proc. Natl. Acad. Sci. U.S.A. 91:12105-12109 (1994).
CC -!- FUNCTION: ACTIVATOR OF YEAST TRANSCRIPTION FACTOR, STE12.
CC -!- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
CC LAMMER SUBFAMILY.
CC -----
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CC -----
CC EMBL; U16176; AAA57117.1; -.

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DR HSP; P24941; 1A01.
DR InterPro; IPR000719; Euk_pkinase.
DR InterPro; IPR002290; Ser_thr_pkinase.
DR Pfam; PF00669; pkinase; 1.
DR ProDom; PD000001; Euk_pkinase; 1.
DR SMART; SM00220; S_TKG; 1.
DR PROSITE; PS00107; PROTEIN KINASE_ATP; FALSE_NEG.
DR PROSITE; PS00108; PROTEIN KINASE_ST; 1.
DR PROSITE; PS50011; PROTEIN KINASE_DOM; 1.
KW Transferase; Serine/threonine-protein kinase; ATP-binding.
FT DOMAIN 115 443 PROTEIN KINASE.
FT NP_BIND 121 129 ATP (BY SIMILARITY).
FT BINDING 144 144 ATP (BY SIMILARITY).
FT ACT_SITE 240 240 BY SIMILARITY.
SQ SEQUENCE 467 AA; 54216 MW; 54D739A82F490E12 CRC64;

Query Match 57.8%; Score 52; DB 1; Length 467;
Best Local Similarity 52.4%; Pred. NO. 15;
Matches 11; Conservative 1; Mismatches 3; Indels 6; Gaps 1;

QY 1 RRRPR-----PPVLPRLPP 15
|:|:|:|:|:|
DB 35 RKRPLTWDAAPLPPPPPP 55

RESULT 13
RELA_STRAT
ID RELA_STRAT STANDARD; PRT; 841 AA.
AC O85709;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE GTP pyrophosphokinase (EC 2.7.6.5) (ATP:GTP 3'-pyrophosphotransferase)
DE (PPGPP synthetase I) (PPGPP synthetase).
GN RELA.
OS Streptomyces antibioticus.
OC Bacteria; Actinobacteria; Actinobacteria (class); Actinobacteridae;
OC Actinomycetales; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=1890;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IMRU 3720;
RX MEDLINE=99296594; PubMed=10368159;
RA Hoyt S., Jones G.H.;
RT "relA is required for actinomycin production in Streptomyces
RT antibioticus.";
RL J. Bacteriol. 181:3824-3829 (1999).
CC -!- FUNCTION: In eubacteria PPGPP (guanosine 3'-diphosphate 5'-
CC diphosphate) is a mediator of the stringent response that
CC coordinates a variety of cellular activities in response to
CC changes in nutritional abundance. This enzyme catalyzes the
CC formation of PPGPP which is then hydrolysed to form PPGPP (By
CC similarity). Is required for actinomycin production.
CC -!- CATALYTIC ACTIVITY: ATP + GTP = AMP + guanosine 3'-diphosphate 5'-
CC triphosphate.
CC -!- PATHWAY: FIRST STEP IN THE METABOLISM OF PPGPP.
CC -!- SIMILARITY: BELONGS TO THE RELA / SPOT FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AF072829; AAC26021.1; -.
CC DR InterPro; IPR002912; ACT.
CC DR InterPro; IPR002819; HD.
CC DR InterPro; IPR003607; ME_Ppase_HDC.
CC DR InterPro; IPR004811; Spot_rela.
CC DR InterPro; IPR004095; TGS_dom.
CC DR Pfam; PF01842; ACT; 1.

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DR Pfam; PF01966; HD; 1.
DR Pfam; PF02824; TGS; 1.
DR SMART; SM00471; HDC; 1.
DR TIGRFAMS; TIGR00691; spot_rela; 1.
DR TRANSFERASE; Kinase.
SQ SEQUENCE 841 AA; 93671 MW; 632A037BA4BF4C94 CRC64;

Query Match 57.8%; Score 52; DB 1; Length 841;
Best Local Similarity 60.0%; Pred. No. 26;
Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 1 RRRPPYLPRLPRPP 15
   |||||
Db 50 RPKPAPPRPPPP 64

RESULT 14
PTNE HUMAN STANDARD; PRT; 1187 AA.
AC Q15678;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Protein tyrosine phosphatase, non-receptor type 14 (EC 3.1.3.48)
DE (Protein-tyrosine phosphatase pez).
GN PTPN14 OR PEZ.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Breast carcinoma;
RX MEDLINE=95251727; PubMed=7733990;
RA Smith A.L., Mitchell P.J., Shipley J., Gusterson B.A., Rogers M.V.,
RA Crompton M.R.;
RT "Pez: a novel human cDNA encoding protein tyrosine phosphatase- and
RT ezrin-like domains.";
RL Biochem. Biophys. Res. Commun. 209:959-965(1995).
CC -1- CATALYTIC ACTIVITY: Protein tyrosine phosphatase + H(2)O = protein
CC tyrosine + phosphate.
CC -1- TISSUE SPECIFICITY: EXPRESSED IN A VARIETY OF HUMAN TISSUES
CC INCLUDING KIDNEY, SKELETAL MUSCLE, LUNG AND PLACENTA.
CC -1- SIMILARITY: CONTAINS 1 BAND 4.1-LIKE DOMAIN.
CC -1- SIMILARITY: BELONGS TO THE NON-RECEPTOR CLASS OF THE PROTEIN-
CC TYROSINE PHOSPHATASE FAMILY.
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EMBL; X82676; CAA57993.1; -
DR HSSP; P29350; LGWZ.
DR Genew; HGNC:9647; PTPN14.
DR MIM; 603155; -
DR InterPro; IPR000299; Band 4.1.
DR InterPro; IPR000387; TYR_PTPase.
DR InterPro; IPR000242; TYR_PP.
DR Pfam; PF00102; Y_phosphatase; 1.
DR Pfam; PF00373; Band_41; 1.
DR PRINTS; PR00935; BAND41.
DR PRINTS; PR00700; PRTYPHPTASE.
DR SMART; SM00194; PTPC; 1.
DR SMART; SM00295; B41; 1.
DR PROSITE; PS00660; BAND_41_1; 1.
DR PROSITE; PS00661; BAND_41_2; 1.
DR PROSITE; PS00657; BAND_41_3; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS00055; TYR_PHOSPHATASE_PTP; 1.

DR PROSITE; PS50056; TYR_PHOSPHATASE_2; 1.
DR Structural protein; Cytoskeleton; Hydrolase.
DR DOMAIN 25 1187 BAND 4.1-LIKE.
DR DOMAIN 933 1187 PROTEIN-TYROSINE PHOSPHATASE.
DR ACT_SITE 1121 1121 BY SIMILARITY.
DR FT DOMAIN 566 573 POLY-PRO.
DR FT DOMAIN 709 716 POLY-GLU.
SQ SEQUENCE 1187 AA; 135239 MW; 015760B75E3574E3 CRC64;

Query Match 57.2%; Score 51.5; DB 1; Length 1187;
Best Local Similarity 83.3%; Pred. No. 42;
Matches 10; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 3 RRRPPYLPRLPRPP 14
   |||||
Db 565 RPPPPY-PRPRP 575

RESULT 15
PTNE MOUSE STANDARD; PRT; 1189 AA.
AC Q62130;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Protein tyrosine phosphatase, non-receptor type 14 (EC 3.1.3.48)
DE (Protein-tyrosine phosphatase PTP36).
GN PTPN14.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CB-17-SCID; TISSUE=Thymus;
RX MEDLINE=94354845; PubMed=8074693;
RA Sawada M., Ogata M., Fujino Y., Hamaoka T.;
RT "cDNA cloning of a novel protein tyrosine phosphatase with homology
RT to cytoskeletal protein 4.1 and its expression in T-lineage cells.";
RL Biochem. Biophys. Res. Commun. 203:479-484(1994).
CC -1- FUNCTION: MAY BE INVOLVED IN THE REGULATION OF T CELL DEVELOPMENT.
CC -1- CATALYTIC ACTIVITY: Protein tyrosine phosphatase + H(2)O = protein
CC tyrosine + phosphate.
CC -1- TISSUE SPECIFICITY: THYMUS; IN CELLS OF BOTH HEMATOPOIETIC AND
CC NON-HEMATOPOIETIC ORIGINS.
CC -1- SIMILARITY: CONTAINS 1 BAND 4.1-LIKE DOMAIN.
CC -1- SIMILARITY: BELONGS TO THE NON-RECEPTOR CLASS OF THE PROTEIN-
CC TYROSINE PHOSPHATASE FAMILY.
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EMBL; D31842; BAA06628.1; -
DR HSSP; Q06124; ZSHP.
DR MGD; MGI:102467; Ptpn14.
DR InterPro; IPR000299; Band 4.1.
DR InterPro; IPR000387; TYR_PTPase.
DR InterPro; IPR000242; TYR_PP.
DR Pfam; PF00102; Y_phosphatase; 1.
DR Pfam; PF00373; Band_41; 1.
DR PRINTS; PR00935; BAND41.
DR PRINTS; PR00700; PRTYPHPTASE.
DR SMART; SM00194; PTPC; 1.
DR SMART; SM00295; B41; 1.
DR PROSITE; PS00660; BAND_41_1; 1.
DR PROSITE; PS00661; BAND_41_2; 1.
DR PROSITE; PS00657; BAND_41_3; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS00055; TYR_PHOSPHATASE_PTP; 1.

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DR PROSITE; PS50055; TYR\_PHOSPHATASE\_PTP; 1.  
DR PROSITE; PS50056; TYR\_PHOSPHATASE\_2; 1.  
KW Structural protein; Cytoskeleton; Hydrolase.  
FT DOMAIN 75 239 BAND 4.1-LIKE.  
FT DOMAIN 935 1189 PROTEIN-TYROSINE PHOSPHATASE.  
FT ACT SITE 1123 1123 BY SIMILARITY.  
FT DOMAIN 566 573 POLY-PRO.  
FT DOMAIN 635 639 POLY-GLY.  
FT DOMAIN 712 718 POLY-GLU.  
SQ SEQUENCE 1189 AA; 135030 MW; 2B85BE5F9C723303 CRC64;  
  
Query Match 57.2%; Score 51.5; DB 1; Length 1189;  
Best Local Similarity 83.3%; Pred. NO. 42;  
Matches 10; Conservative 0; Mismatches 1; Indels 1; Gaps 1;  
  
QY 3 RPPPPYLP RPP 14  
DB 565 RPPPPY-P RPP 575

Search completed: May 13, 2003, 10:40:51  
Job time : 13 secs

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GenCore version 5.1.4.p5.4578  
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OM protein - protein search, using sw model

Run on: May 13, 2003, 10:37:02 ; Search time 29 seconds  
(without alignments)  
106.576 Million cell updates/sec

Title: US-09-426-011d-3

Perfect score: 90

Sequence: 1 RRRPPVLPVLRPP 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 671580 seqs, 206047115 residues

Total number of hits satisfying chosen parameters: 671580

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL\_21.\*

1: sp\_archaea.\*

2: sp\_bacteria.\*

3: sp\_fungi.\*

4: sp\_human.\*

5: sp\_invertebrate.\*

6: sp\_mammal.\*

7: sp\_mhc.\*

8: sp\_organelle.\*

9: sp\_phase.\*

10: sp\_plant.\*

11: sp\_rodent.\*

12: sp\_virus.\*

13: sp\_vertebrate.\*

14: sp\_unclassified.\*

15: sp\_rvirus.\*

16: sp\_bacteriaph.\*

17: sp\_archaeap.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	61	67.8	336	12 Q68405	Q68405 human cytom
2	58	64.4	156	10 Q8RV32	Q8RV32 oryza sativ
3	58	64.4	164	6 P79361	P79361 ovis aries
4	58	64.4	190	6 Q9X3Q9	Q9X3Q9 capra hircu
5	58	64.4	224	6 O19031	O19031 ovis aries
6	57.5	63.9	183	10 Q94J79	Q94J79 oryza sativ
7	57	63.3	200	16 Q9RK54	Q9RK54 streptomyce
8	57	63.3	361	2 Q9XCG4	Q9XCG4 mycobacteri
9	56	62.2	212	2 O08306	O08306 nocardioid
10	55	61.1	212	12 O41980	O41980 murid herpe
11	55	61.1	464	12 Q91TM2	Q91TM2 tupaia herp
12	54.5	60.6	301	10 Q41848	Q41848 zea mays (m
13	54.5	60.6	2635	12 P88955	P88955 kaposi's sa
14	54.5	60.6	2635	12 O40942	O40942 kaposi's sa
15	54	60.0	347	11 Q8R353	Q8R353 mus musculu
16	54	60.0	359	5 Q9XZT0	Q9XZT0 drosophila

17	54	60.0	427	5	O44582	O44582 caenorhabdi
18	54	60.0	451	5	Q95X63	Q95X63 caenorhabdi
19	54	60.0	955	4	Q9Y2W1	Q9Y2W1 homo sapien
20	53.5	59.4	225	11	Q99JA6	Q99JA6 mus musculu
21	53	58.9	168	10	Q9SM77	Q9SM77 oryza sativ
22	53	58.9	184	5	Q23291	Q23291 caenorhabdi
23	53	58.9	185	10	Q94JF6	Q94JF6 oryza sativ
24	53	58.9	199	5	Q8WSY8	Q8WSY8 apis mellif
25	53	58.9	333	10	Q9X123	Q9X123 oryza sativ
26	53	58.9	428	10	Q23370	Q23370 arabidopsis
27	53	58.9	491	10	O82066	O82066 solanum tub
28	53	58.9	520	10	Q9LV14	Q9LV14 arabidopsis
29	52.5	58.3	602	12	Q66852	Q66852 fowl adenov
30	52	57.8	148	16	Q8U5T2	Q8U5T2 agrobacteri
31	52	57.8	155	4	Q96E55	Q96E55 homo sapien
32	52	57.8	439	10	Q42421	Q42421 beta vulgar
33	52	57.8	450	10	Q94CE1	Q94CE1 arabidopsis
34	52	57.8	467	10	Q39184	Q39184 arabidopsis
35	52	57.8	790	5	Q8T458	Q8T458 drosophila
36	52	57.8	1006	10	Q9LMQ1	Q9LMQ1 arabidopsis
37	52	57.8	1091	5	Q9W1Z6	Q9W1Z6 drosophila
38	51.5	57.2	145	12	Q8V718	Q8V718 simian herp
39	51.5	57.2	238	10	Q8W097	Q8W097 oryza sativ
40	51	56.7	94	5	Q917F1	Q917F1 drosophila
41	51	56.7	145	10	Q8SAY8	Q8SAY8 oryza sativ
42	51	56.7	255	10	Q8RYX5	Q8RYX5 oryza sativ
43	51	56.7	409	5	Q9U0Z7	Q9U0Z7 leishmania
44	51	56.7	417	5	Q9V4V1	Q9V4V1 drosophila
45	51	56.7	470	10	Q8S5U3	Q8S5U3 oryza sativ

#### ALIGNMENTS

#### RESULT 1

Q68405 PRELIMINARY; PRT; 336 AA.  
AC Q68405;  
DT 01-NOV-1996 (T-EMBLrel. 01, Created)  
DT 01-NOV-1996 (T-EMBLrel. 01, Last sequence update)  
DT 01-NOV-1998 (T-EMBLrel. 08, Last annotation update)  
DE Orf UL151.  
OS Human cytomegalovirus.  
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
OC Betaherpesvirinae; Cytomegalovirus.  
OX NCBI\_TaxID=10359;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=TOLEDO;  
RX MEDLINE=96099416; PubMed=8523595;  
RA Cha T.A., Tom E., Kemble G.W., Duke G.M., Mocarski E.S., Spaete R.R.;  
RT "Human cytomegalovirus clinical isolates carry at least 19 genes not  
found in laboratory strains.";  
RL J. Virol. 70:78-83(1996).  
DR EMBL; U33331; AAA8582.1; --  
SQ SEQUENCE 336 AA; 35116 MW; 9F865E5019F69D0C CRC64;

Query Match 67.8%; Score 61; DB 12; Length 336;  
Best Local Similarity 78.6%; Pred. No. 0.94;  
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 RRRPPVLPVLRPP 15

Db 279 RRRPPIQLQRP 292

#### RESULT 2

Q8RV32 PRELIMINARY; PRT; 156 AA.

AC Q8RV32;

DT 01-JUN-2002 (T-EMBLrel. 21, Created)

DT 01-JUN-2002 (T-EMBLrel. 21, Last sequence update)

DT 01-JUN-2002 (T-EMBLrel. 21, Last annotation update)

DE OSJNB0032K15.1 protein (OJ1159.D09.32 protein).  
GN OSJNB0032K15.1 OR OJ1159.D09.32.  
OS Oryza sativa (japonica cultivar-group).  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
OC Ehrhartoideae; Oryzeae; Oryza.  
OX NCBI\_TaxID=39947;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CV. NIPPONBARE;  
RA Sasaki T., Matsumoto T., Yamamoto K.;  
RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC  
clone:OSJNB0032K15."  
RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.  
RN [2]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CV. NIPPONBARE;  
RA Sasaki T., Matsumoto T., Yamamoto K.;  
RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC  
clone:OJ1159.D09."  
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
DR EMBL; AP003710; BAB86560.1;  
DR EMBL; AP003792; BAB89214.1;  
SQ SEQUENCE 156 AA; 17659 MW; 4152112C3DB493CF CRC64;  
Query Match 64.4%; Score 58; DB 10; Length 156;  
Best Local Similarity 73.3%; Pred. No. 1.2; Mismatches 0; Indels 0; Gaps 0;  
Matches 11; Conservative 0;  
QY 1 RRRPRPPYLP RPRP 15  
DB 78 RRRPRPPYLP RPRP 92  
RESULT 3  
ID P79361 PRELIMINARY; PRT; 164 AA.  
AC P79361;  
DT 01-MAY-1997 (TrEMBLrel. 03, Created)  
DT 01-MAY-1997 (TrEMBLrel. 03, Last sequence update)  
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)  
DE 7.5 kDa bactinecin (Fragment).  
GN BAC7.5.  
OS Ovis aries (Sheep).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
OC Bovidae; Caprinae; Ovis.  
OX NCBI\_TaxID=9940;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC TISSUE=LIVER;  
RX MEDLINE=96140581; PubMed=8549789;  
RA Mahoney M.M., Lee A.Y., Brezinski-Caliguri D.J., Huttner K.M.;  
RT "Molecular analysis of the sheep cathelin family reveals a novel  
antimicrobial peptide."  
RL FEBS Lett. 377:519-522(1995).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC TISSUE=LIVER;  
RA Huttner K.M., Mahoney M.M.;  
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.  
DR EMBL; U60598; AAB49713.1;  
DR InterPro; IPR001894; Cathelicidin.  
DR Pfam; PF00666; Cathelicidins; 1.  
DR ProDom; PD001838; Cathelicidin; 1.  
DR PROSITE; PS00946; CATHELICIDINS\_1; 1.  
DR PROSITE; PS00947; CATHELICIDINS\_2; 1.  
FT NON TER 164  
SQ SEQUENCE 164 AA; 18642 MW; E3BFC871F6AE8B9A CRC64;  
Query Match 64.4%; Score 58; DB 6; Length 164;  
Best Local Similarity 78.6%; Pred. No. 1.2; Mismatches 0; Indels 0; Gaps 0;  
Matches 11; Conservative 0;

QY 1 RRRPRPPYLP RPRP 14  
DB 132 RLRPRRPLRPRP 145  
RESULT 4  
O9XSQ9  
ID O9XSQ9 PRELIMINARY; PRT; 190 AA.  
AC O9XSQ9;  
DT 01-NOV-1999 (TrEMBLrel. 12, Created)  
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)  
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)  
DE BAC7.5 protein.  
GN BAC7.5.  
OS Capra hircus (Goat).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
OC Bovidae; Caprinae; Capra.  
OX NCBI\_TaxID=9925;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC TISSUE=BONE MARROW;  
RA Zhao C., Nguyen T., Brogden K., Lehrer R.;  
RT "cDNA cloning of goat cathelin related peptides."  
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.  
DR EMBL; AJ243125; CAB45523.1;  
DR InterPro; IPR001894; Cathelicidin.  
DR Pfam; PF00666; Cathelicidins; 1.  
DR ProDom; PD001838; Cathelicidin; 1.  
DR PROSITE; PS00946; CATHELICIDINS\_1; 1.  
DR PROSITE; PS00947; CATHELICIDINS\_2; 1.  
FT CHAIN 131  
SQ SEQUENCE 190 AA; 21835 MW; D13305EF16875F4F CRC64;  
Query Match 64.4%; Score 58; DB 6; Length 190;  
Best Local Similarity 78.6%; Pred. No. 1.4; Mismatches 0; Indels 0; Gaps 0;  
Matches 11; Conservative 0;  
QY 1 RRRPRPPYLP RPRP 14  
DB 132 RLRPRRPLRPRP 145  
RESULT 5  
O19031  
ID O19031 PRELIMINARY; PRT; 224 AA.  
AC O19031;  
DT 01-JAN-1998 (TrEMBLrel. 05, Created)  
DT 01-JAN-1998 (TrEMBLrel. 05, Last sequence update)  
DT 01-JUN-2001 (TrEMBLrel. 17, Last annotation update)  
DE BACTINECIN 11 precursor.  
GN BAC11.  
OS Ovis aries (Sheep).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
OC Bovidae; Caprinae; Ovis.  
OX NCBI\_TaxID=9940;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC MEDLINE=98121317; PubMed=9461419;  
RA Huttner K.M., Lambeth M.R., Burkin H.R., Broad T.E.;  
RT "Localization and genomic organization of sheep antimicrobial peptides  
genes."  
RL Gene 206:85-91(1998).  
CC -!- FUNCTION: ANTIMICROBIAL PEPTIDE (BY SIMILARITY).  
CC -!- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.  
DR EMBL; U77049; AAB62000.1;  
DR EMBL; U77046; AAB62000.1; JOINED.  
DR EMBL; U77047; AAB62000.1; JOINED.  
DR EMBL; U77048; AAB62000.1; JOINED.  
DR InterPro; IPR001894; Cathelicidin.  
DR Pfam; PF00666; Cathelicidins; 1.

DR ProDom; PD001838; Cathelicidin; 1.  
DR PROSITE; PS00946; CATHELICIDINS 1; 1.  
DR PROSITE; PS00947; CATHELICIDINS 2; 1.  
KW Signal; Antibiotic.  
FT SIGNAL 1 29 POTENTIAL.  
FT PROPEP 30 130 POTENTIAL.  
FT CHAIN 131 224 BACTINECIN 11.  
FT MOD\_RES 30 30 PYROLIDONE CARBOXYLIC ACID (BY  
FT SIMILARITY).  
FT DISULFID 85 96 BY SIMILARITY.  
FT DISULFID 107 124 BY SIMILARITY.  
SQ SEQUENCE 224 AA; 25669 MW; 6AEAB1256AC76FC CRC64;  
Query Match 64.4%; Score 58; DB 6; Length 224;  
Best Local Similarity 78.6%; Pred. No. 1.6;  
Matches 11; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
QY 1 RRRPRPPYLPRLPR 14  
DB 132 RLRRPRRLPRPP 145  
RESULT 6  
Q94J98 PRELIMINARY; PRT; 183 AA.  
AC Q94J98;  
DT 01-DEC-2001 (TrEMBLrel. 19, Created)  
DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)  
DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)  
DE P0047B08.14 protein (OJ1159.D09.5 protein).  
GN P0047B08.14 OR OJ1159.D09.5.  
OS Oryza sativa (Rice), and  
OS Oryza sativa (japonica cultivar-group).  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;  
OC Ehrhartoideae; Oryzaceae; Oryza.  
OX NCBI\_TaxID=4530, 39947;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CV. NIPPONBARE;  
RA Sasaki T., Matsumoto T., Yamamoto K.;  
RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, PAC  
clone: P0047B08.";  
RL Submitted (DEC-2000) to the EMBL/GenBank/DBJ databases.  
RN [2]  
RP SEQUENCE FROM N.A.  
RC STRAIN=CV. NIPPONBARE;  
RA Sasaki T., Matsumoto T., Yamamoto K.;  
RT "Oryza sativa (japonica cultivar-group) genomic DNA, chromosome 1, BAC  
clone: OJ1159.D09.";  
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
DR EMBL; AP003053; BAB55690.1;  
DR EMBL; AP003792; BAB89188.1;  
SQ SEQUENCE 183 AA; 20155 MW; F1CF82AD89CEB36 CRC64;  
Query Match 63.9%; Score 57.5; DB 10; Length 183;  
Best Local Similarity 73.3%; Pred. No. 1.6;  
Matches 11; Conservative 1; Mismatches 2; Indels 1; Gaps 1;  
QY 1 RRRPRPPYLPRLPRPP 15  
DB 129 RSRPR-PYAPRPQP 142  
RESULT 7  
Q9RK54 PRELIMINARY; PRT; 200 AA.  
ID Q9RK54  
AC Q9RK54;  
DT 01-MAY-2000 (TrEMBLrel. 13, Created)  
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)  
DT 01-JUN-2002 (TrEMBLrel. 21, Last annotation update)  
DE Hypothetical protein SCO0323.  
GN SCO0323 OR SCF12.02C.

OS Streptomyces coelicolor.  
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;  
OC Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.  
OX NCBI\_TaxID=1902;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=A3(2) / M145;  
RA Bentley S.D., Chater K.F., Cerdano-Tarraga A.-M., Challis G.L.,  
Thomson N.R., James K.D., Harris D.E., Quail M.A., Kieser H.,  
Harper D., Bateman A., Brown S., Chandra G., Chen C.W., Collins M.,  
Cronin A., Fraser A., Goble T., Larke L., Murphy L., Oliver K., O'Neill S.,  
Huang C.-H., Kieser T., Lark M., Rutherford K., Rutter S.,  
Rabinowitz E., Rajandream M.A., Rutherford K., Rutter S.,  
Seeger K., Saunders D., Sharp S., Squares R., Taylor K.,  
Warren T., Wietzorrek A., Woodward J., Barrell B.G., Parkhill J.,  
Hopwood D.A.;  
RT "Complete genome sequence of the model actinomycete Streptomyces  
coelicolor A3(2).";  
RL Nature 417:141-147(2002).  
DR EMBL; AL117669; CAB56128.1; --  
KW Hypothetical protein.  
SQ SEQUENCE 200 AA; 22076 MW; 0DCBBEC5585803B5 CRC64;  
Query Match 63.3%; Score 57; DB 16; Length 200;  
Best Local Similarity 76.9%; Pred. No. 2;  
Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
QY 1 RRRPRPPYLPRLPR 13  
DB 118 RRHPEPPALPRPP 130  
RESULT 8  
Q9XCG4 PRELIMINARY; PRT; 361 AA.  
ID Q9XCG4;  
AC Q9XCG4;  
DT 01-NOV-1999 (TrEMBLrel. 12, Created)  
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)  
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)  
DE Hypothetical 40.2 kDa protein.  
OS Mycobacterium avium.  
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;  
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.  
OX NCBI\_TaxID=1764;  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=2151;  
RA Eckstein T.M., Lambert M.L., Brennan P.J., Belisle J.T., Inamine J.M.;  
RT "Identification of a gene cluster involved in glycopeptidolipid  
biosynthesis and of a gene cluster encoding daunorubicin resistance in  
two strains of Mycobacterium avium serovar 2.";  
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.  
DR EMBL; AF143772; AAD44199.1; --  
KW Hypothetical protein.  
SQ SEQUENCE 361 AA; 40208 MW; AD01DBE825C1C9EA CRC64;  
Query Match 63.3%; Score 57; DB 2; Length 361;  
Best Local Similarity 71.4%; Pred. No. 3.4;  
Matches 10; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
QY 1 RRRPRPPYLPRLPRPP 14  
DB 32 RRRPRPPAPHPPP 45  
RESULT 9  
Q08306 PRELIMINARY; PRT; 212 AA.  
ID Q08306  
AC Q08306;  
DT 01-JUL-1997 (TrEMBLrel. 04, Created)  
DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)  
DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)  
DE Hypothetical 22.7 kDa protein.

OS Nocardioideis simplex (Arthrobacter simplex).  
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;  
 OC Actinomycetales; Propionibacterineae; Nocardioideaceae; Pimelobacter.  
 OX NCBI\_TaxID=2045;  
 RN [1]  
 RP SEQUENCE FROM N.A.,  
 RC STRAIN=IFO12069;  
 RX MEDLINE=95319331; PubMed=7596291;  
 RA Molnar I., Choi K., Yamashita M., Murooka Y.;  
 RT "Molecular cloning, expression in Streptomyces lividans, and analysis  
 of a gene cluster from Arthrobacter simplex encoding 3-  
 ketosteroid-DELTA.1.1-dehydrogenase, 3-ketosteroid-DELTA.5-isomerase  
 RT and a hypothetical regulatory protein.";  
 RL Mol. Microbiol. 15:895-905(1995).  
 RN [2]  
 RP SEQUENCE FROM N.A.,  
 RC STRAIN=IFO12069;  
 RX MEDLINE=95319331; PubMed=7596291;  
 RA Dzialek J., Yamashita M., Murooka Y.;  
 RT "Cloning, sequencing and characterization of the downstream region of  
 the kdsI operon of Arthrobacter simplex.";  
 RL Submitted (MAR-1997) to the EMBL/GenBank/DBJ databases.  
 CC -1- SIMILARITY: BELONGS TO THE TETR/ACRR FAMILY OF TRANSCRIPTIONAL  
 REGULATORS.  
 DR EMBL; Z93338; CAB07542.1; -;  
 DR InterPro; IPR001647; HTH\_Tetr.  
 DR Pfam; PF00440; tetr; 1.  
 DR PRINTS; PR00455; HTHTETR.  
 DR PROSITE; PS01081; HTH\_TETR\_FAMILY; 1.  
 KW DNA-binding; Hypothetical protein; Transcription regulation.  
 SQ SEQUENCE 212 AA; 22740 MW; F9118E18DDF4E0B2 CRC64;

Query Match 62.2%; Score 56; DB 2; Length 212;  
 Best Local Similarity 73.3%; Pred. No. 2.8;  
 Matches 11; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRRPPYLPFRPP 15  
 DB 83 RRRPPGSGRPP 97

RESULT 10

O41980  
 ID O41980 PRELIMINARY; PRT; 212 AA.  
 AC O41980;  
 DT 01-JAN-1998 (T-Emblrel. 05, Created)  
 DT 01-JAN-1998 (T-Emblrel. 05, Last sequence update)  
 DT 01-JUN-2000 (T-Emblrel. 14, Last annotation update)  
 DE Hypothetical 21.9 kDa protein.  
 GN GAMMAHV.M13.  
 OS murid herpesvirus 4.  
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Gammaherpesvirinae.  
 OX NCBI\_TaxID=33708;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=WUMS;  
 RX MEDLINE=97366649; PubMed=9223479;  
 RA Virgin H.W. IV, Latrelle P., Wamsley P., Hallsworth K., Weck K.E.,  
 Dal Canto A.J., Speck S.H.;  
 RT "Complete sequence and genomic analysis of murine gammaherpesvirus  
 68.";  
 RL J. Virol. 71:5894-5904(1997).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=WUMS;  
 RA Latrelle P., Wamsley P., Waterston R.H.;  
 RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; U97553; AAB66426.1; -;  
 KW Hypothetical protein.  
 SQ SEQUENCE 212 AA; 21911 MW; E066860064282149 CRC64;

Query Match 61.1%; Score 55; DB 12; Length 212;  
 Best Local Similarity 75.0%; Pred. No. 3.8;

Matches 9; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 QY 4 PRPPYLPFRPP 15  
 DB 136 PSPPPLPRQPP 147

RESULT 11

O91TM2  
 ID O91TM2 PRELIMINARY; PRT; 464 AA.  
 AC O91TM2;  
 DT 01-DEC-2001 (T-Emblrel. 19, Created)  
 DT 01-DEC-2001 (T-Emblrel. 19, Last sequence update)  
 DT 01-DEC-2001 (T-Emblrel. 19, Last annotation update)  
 DE T74.  
 OS Tupaia herpesvirus.  
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Betaherpesvirinae.  
 OX NCBI\_TaxID=10397;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=2;  
 RX MEDLINE=21211637; PubMed=11312357;  
 RA Bahr U., Darai G.;  
 RT "Analysis and Characterization of the Complete Genome of Tupaia (Tree  
 Shrew) Herpesvirus.";  
 RL J. Virol. 75:4854-4870(2001).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=2;  
 RA Darai G., Bahr U.;  
 RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; AF281817; AAKS7119.1; -;  
 SQ SEQUENCE 464 AA; 51193 MW; 4BB7313EA2C2BD16 CRC64;

Query Match 61.1%; Score 55; DB 12; Length 464;  
 Best Local Similarity 76.9%; Pred. No. 7.7;  
 Matches 10; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 RRRPPYLPFRPP 15  
 DB 421 RRRPPRPRPP 433

RESULT 12

O41848  
 ID O41848 PRELIMINARY; PRT; 301 AA.  
 AC O41848;  
 DT 01-NOV-1996 (T-Emblrel. 01, Created)  
 DT 01-NOV-1996 (T-Emblrel. 01, Last sequence update)  
 DT 01-JUN-2002 (T-Emblrel. 21, Last annotation update)  
 DE Prolin rich protein.  
 GN PRP.  
 OS Zea mays (Maize).  
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC clade;  
 OC Panicoideae; Andropogoneae; Zea.  
 OX NCBI\_TaxID=4577;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=W64A;  
 RX MEDLINE=92361259; PubMed=1498600;  
 RA Jose-Estanyol M., Ruiz-Avila L., Puigdomenech P.;  
 RT "A maize embryo-specific gene encodes a proline-rich and hydrophobic  
 protein.";  
 RL Plant Cell 4:413-423(1992).  
 DR EMBL; X60432; CAA42959.1; -;  
 DR HSP; P24337; IHVP.  
 DR InterPro; IPR003612; AAI.  
 DR InterPro; IPR001768; Try/amy1 inhbr.  
 DR Pfam; PF00234; try\_alpha\_amy1; 1.  
 DR SMART; SM00499; AAI; 1.  
 SQ SEQUENCE 301 AA; 31647 MW; 884EB70854D28C2E CRC64;

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Query Match          60.6%; Score 54.5; DB 10; Length 301;
Best Local Similarity 71.4%; Pred. No. 6;
Matches 10; Conservative 1; Mismatches 2; Indels 1; Gaps 1;

QY 3 RRPYPYL-PPRPP 15
   |||: |||
Db 149 RSPPPYVPTDRPP 162

RESULT 13
P88955 ID P88955 PRELIMINARY; PRT; 2635 AA.
AC P88955;
DT 01-MAY-1997 (TREMELrel. 03, Created)
DT 01-MAY-1997 (TREMELrel. 03, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE ORF 64.
OS Kaposi's sarcoma-associated herpesvirus (KSHV) (Human herpesvirus 8).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae; Rhadinovirus.
OX NCBI_TaxID=37296;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97094384; PubMed=8939871;
RA Moore P.S., Boshoff C., Weiss R.A., Chang Y.;
RT "Molecular mimicry of human cytokine and cytokine response pathway
   genes by KSHV";
RL Science 274:1739-1744 (1996).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=97121480; PubMed=8962146;
RA Russo J.J., Bohenzky R.A., Chien M.-C., Chen J., Yan M., Maddalena D.,
   Parry J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RA Party J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RT "Nucleotide sequence of the Kaposi sarcoma-associated herpesvirus
   (HHV8)";
RL Proc. Natl. Acad. Sci. U.S.A. 93:14862-14867 (1996).
RN [3]
RP SEQUENCE FROM N.A.
RA Russo J.J., Bohenzky R.A., Chien M.-C., Chen J., Yan M., Maddalena D.,
   Parry J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RA Party J.P., Peruzzi D., Edelman I.S., Chang Y., Moore P.S.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; U75698; AAC57149.1; -.
SQ SEQUENCE 2635 AA; 289687 MW; 00070132EA8139AF CRC64;

Query Match          60.6%; Score 54.5; DB 12; Length 2635;
Best Local Similarity 68.4%; Pred. No. 44;
Matches 13; Conservative 0; Mismatches 1; Indels 5; Gaps 2;

QY 2 RRPR---PPYLP-RRPRP 15
   |||: |||
Db 271 RRPVVIPYDPTDRPP 289

RESULT 14
O40942 ID O40942 PRELIMINARY; PRT; 2635 AA.
AC O40942;
DT 01-JAN-1998 (TREMELrel. 05, Created)
DT 01-JAN-1998 (TREMELrel. 05, Last sequence update)
DT 01-DEC-2001 (TREMELrel. 19, Last annotation update)
DE ORF 64.
OS Kaposi's sarcoma-associated herpesvirus (KSHV) (Human herpesvirus 8).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Gammaherpesvirinae; Rhadinovirus.
OX NCBI_TaxID=37296;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97296220; PubMed=9151804;
RA Neipel F., Albrecht J.C., Fleckenstein B.;
RT "Cell-homologous genes in the Kaposi's sarcoma-associated rhadinovirus
   human herpesvirus 8: determinants of its pathogenicity?";
RL J. Virol. 71:4187-4192 (1997).
```

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[2]
RN SEQUENCE FROM N.A.
RP Neipel F., Albrecht J.-C., Ensser A., Huang Y.-Q., Li J.J.,
   Friedman-Kien A.E., Fleckenstein B.;
RT "The genome of human herpesvirus 8 cloned from Kaposi's sarcoma.";
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; U93872; AAB62600.1; -.
SQ SEQUENCE 2635 AA; 289717 MW; 91DDA0D6FF7B660A CRC64;

Query Match          60.6%; Score 54.5; DB 12; Length 2635;
Best Local Similarity 68.4%; Pred. No. 44;
Matches 13; Conservative 0; Mismatches 1; Indels 5; Gaps 2;

QY 2 RRPR---PPYLP-RRPRP 15
   |||: |||
Db 271 RRPVVIPYDPTDRPP 289

RESULT 15
O8R353 ID O8R353 PRELIMINARY; PRT; 347 AA.
AC O8R353;
DT 01-JUN-2002 (TREMELrel. 21, Created)
DT 01-JUN-2002 (TREMELrel. 21, Last sequence update)
DT 01-JUN-2002 (TREMELrel. 21, Last annotation update)
DE Similar to thyroid hormone receptor-associated protein, 150 kDa
   subunit.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Strausberg R.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC026580; AAH26580.1; -.
KW Receptor.
SQ SEQUENCE 347 AA; 38293 MW; C885A3C2394F4DA6 CRC64;

Query Match          60.0%; Score 54; DB 11; Length 347;
Best Local Similarity 60.0%; Pred. No. 8;
Matches 9; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 RRPYPYL-PPRPP 15
   |||: |||
Db 262 RSPPPYVPTDRPP 276

Search completed: May 13, 2003, 10:41:27
Job time : 32 secs
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